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Sampling and Analysis Plan for Removal of Structures External to the 100K Storage Basins

Prepared for the U.S. Department of Energy Assistant Secretary for Environmental Management

Project Hanford Management Contractor for the U.S. Department of Energy under Contract DE-AC06-96RL13200

Fluor Hanford

Richland, Washington

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Sampling and Analysis Plan for Removal of Structures External to the 100K Storage Basins

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Fluor Hanford

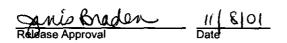
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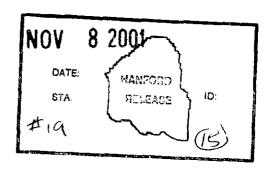
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Sampling and Analysis Plan for Removal of Structures External to the 100 K Storage Basins

October 2001

Prepared by

Environmental Quality Management for Fluor Hanford, Inc.

EXECUTIVE SUMMARY

This Sampling and Analysis Plan (SAP) presents the rationale for waste characterization and the strategy for sampling and analysis activities to support removal of structures and soil external to the K East Fuel Transfer System Building in the K East Area on the Hanford Site. This project is focused on characterization to support waste designation for disposal of the resulting waste at the appropriate disposal facility. This interim remedial action is conducted under the 100 Area Remaining Sites ROD which identifies the Environmental Restoration Disposal Facility (ERDF) as the preferred disposal facility.

The waste that is the subject of this SAP includes:

- Concrete from a pad previously used to store filters from the K East Basin, excluding the highest radiologically contaminated areas,
- Soil underlying the concrete pad and soil on the north side of the pad, down to a depth of between a few inches to 4.5 ft.,
- Sections of the carbon steel monorail and associated concrete supports,
- Asphalt road on the east side of the pad, and
- Transite siding (external wall).

The structures and soil addressed by this SAP are either known to be contaminated with radioactivity or are in areas where contamination is possible. Therefore, all materials removed from the buildings and associated structures are presumed to be radioactively contaminated. All debris and soil will be managed as low-level radioactive waste.

Debris that contains *Resource Conservation and Recovery Act of 1976* (RCRA)/Washington State dangerous constituents above regulated levels will be designated as mixed waste. These constituents may be present at levels that require treatment to comply with Land Disposal Restrictions (LDR). Debris composed primarily of pieces greater than 60 mm, that requires treatment for compliance with the LDR, will be treated through an approved alternative treatment technology for debris under 40 *Code of Federal Regulations* 268.45. Debris less than or equal to 60 mm will be byproducts from and commingled with larger debris items and will be managed with the related waste stream. Only a small amount of debris less than or equal to 60 mm is anticipated.

The sampling design for the debris uses facility or historical radiological sample data to establish the radionuclide distribution of radiological constituents of concern. The radionuclide distributions are established for each waste stream and subsequently used to estimate the content of constituents of concern, indexed to cesium-137 (Cs-137). The Cs-137 content of the waste debris will be estimated using the weight-to-curie relationships previously developed for K Basin above water waste (HNF 2001). Laboratory analysis of the soil underlying and around the filter wash pad will be used to determine the Cs-137 content and the concentration of hazardous constituents. Section 2.3 discusses the details of this approach and utilizes existing sampling and analysis procedures.

Based on operational history, the contamination source term for the concrete, steel, and soil addressed in this SAP is assumed to be the same as the K Basin above water waste (basin water and sludge). Thus, the radionuclide distribution used to estimate the concentration of radionuclides in the waste will be the same as that previously developed for above water debris from K East (HNF 2001).

In cases where assumptions used to establish historical radionuclide ratios are shown not to be applicable, contingency sampling and analysis may be required. Section 2.4 presents methods to obtain contingency laboratory analysis of the debris to measure specific isotopes to revise the radionuclide ratios for a waste stream. Section 2.4 also includes use of nondestructive assay as a contingency analytical approach. It must be emphasized that Section 2.4 is for contingency analysis and not routine use. Fluor Hanford is responsible for contingency sampling.

The concrete pad is painted with different paint found on than the monorail and concrete monorail supports. The material safety data sheets (MSDSs) (Appendix B) for the paint on the pad will be used instead of characterization to develop the waste profile for the pad. No polychlorinated biphenyls are present in this paint. To determine if the concrete debris from the pad designates as RCRA hazardous waste, the concentrations of RCRA constituents in the debris will be based on the concentration of hazardous constituents in the paint being evaluated as a portion of the mass of the debris. Waste smaller than 60 mm will be managed based on a determination of hazardous constituents.

The carbon steel monorail and the concrete foundation and supports were painted with the same paint discussed in HNF 2001, and a previous ERDF profile exists for debris painted with this paint. The same information used for the previous ERDF profile for debris from K Basin will be used to assess the characteristic waste in the monorail debris.

The asphalt roadway is excluded from designation per Washington Administrative Code 173-303-071(3)(e) and no characterization will be done for disposal.

The soil will be sampled to confirm the radionuclide distribution and the concentrations of characteristic metals and organics and additional organics required by ERDF. The details of the sampling are discussed in Section 3.0 of this document.

Personal protective equipment (PPE) associated with these removal activities will be discarded using the same procedures as PPE currently generated during routine K Basins operations. As such, the PPE is covered by other waste profiles currently in place for K Basin PPE. As a result, this SAP does not discuss PPE.

This SAP is based on the results of implementing the Data Quality Objectives (DQO) Process as documented in the Data Quality Objectives Process in Support of the Characterization of Debris and Soil from Removal of Structures External to the 100 K Storage Basins (EQM 2001). The following topics are summarized in Section 1.0:

- historical data,
- rationale for data collection, including radiation surveys and sampling, and
- results of the DQO Process.

Section 2.0 includes the quality assurance project plan that includes details of the radiation survey, analytical methods, detection limits, accuracy, and precision criteria.

Section 3.0 includes the sampling plan that summarizes information needed by those collecting and shipping samples to the laboratory or those performing the radiation surveys.

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ACRONYMS

AJHA automated job hazards analysis ALARA as low as reasonably achievable

BHI Bechtel Hanford, Inc.

CERCLA Comprehensive Environmental Response, Compensation, and Liability Act of

1980

CFR Code of Federal Regulations
CMP Chemical Management Program

COC constituent of concern

COPC constituent of potential concern
DOE U.S. Department of Energy
DQO data quality objectives

DW dangerous waste

EHW extremely hazardous waste

EPA U.S. Environmental Protection Agency environmental restoration contractor

ERDF Environmental Restoration Disposal Facility

FH Fluor Hanford, Inc.
GEA gamma energy analysis
HASP Health and Safety Plan

HASQARD Hanford Analytical Quality Assurance Requirements Documents

HM/HW hazardous material/hazardous waste

HSRCM Hanford Site Radiological Control Manual

ICP inductively coupled plasma (emission spectroscopy)

LDR Land Disposal Restrictions

LLW low-level waste

MSDS material safety data sheet NDA nondestructive assay PCB polychlorinated biphenyl

PHMC Project Hanford Management Contract

PPE personal protective equipment PQL practical quantitation limit

QA quality assurance QC quality control

RCRA Resource Conservation and Recovery Act of 1976

ROD Record of Decision

RPD relative percent difference SAP Sampling and Analysis Plan

SNF spent nuclear fuel

SFL standard fixed laboratory SOP standard operating procedure

TBD to be determined toxicity characteristic

TCLP Toxicity Characteristic Leaching Procedure

TRU transuranic

TSCA	Toxic Substances Control Act of 1976
VOC	volatile organic compound
WAC	waste acceptance criteria

1.0 INTRODUCTION

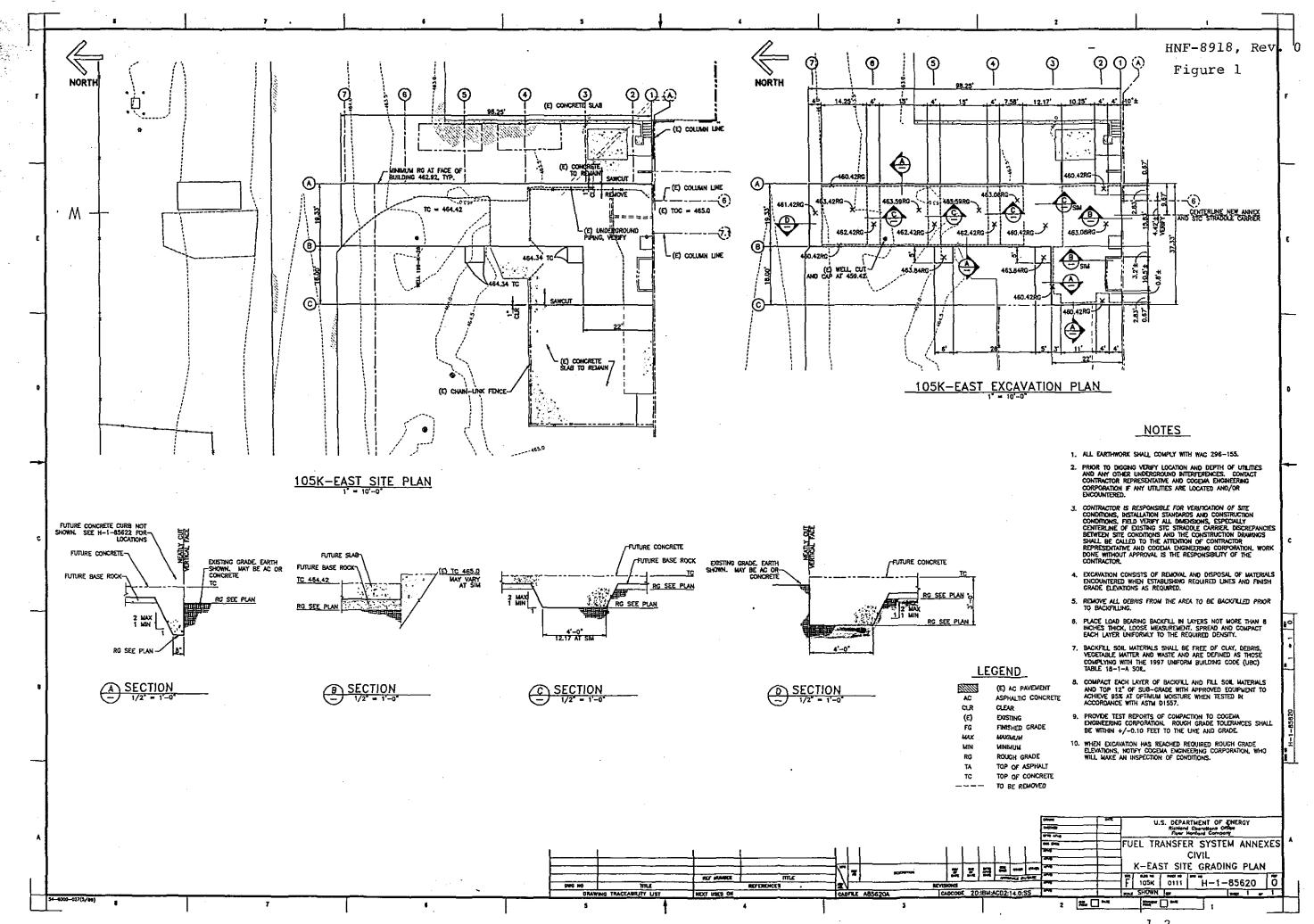
The U.S. Department of Energy (DOE) is currently removing spent nuclear fuel (SNF), sludge, and debris from the K East Area, located in the 100 Area of the Hanford Site. This Sampling and Analysis Plan (SAP) is focused on removal and characterization of the following materials for waste designation and disposal:

- a portion of the filter wash pad,
- the painted and unpainted sections of the carbon steel monorail,
- the painted concrete supports for the monorail,
- the underlying soil,
- the soil that is not covered by any structure on the north side of the pad,
- the asphalt road on the east side of the pad and soil area, and
- the Transite siding (external wall).

All areas are external to the Fuel Transfer System building located in K East.

This work is governed by the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (CERCLA) 100 Areas Remaining Sites ROD (Interim Action Record of Decision for the 100-BC-1, 100-BC-2, 100-DR-1, 100-DR-2, 100-FR-1, 100-FR-2, 100-HR-1, 100-HR-2, 100-KR-1, 100-KR-2, 100-IU-2, 100-IU-6, and 200-CW-3 Operable Units, Hanford Site, Benton County, Washington [EPA 1999]). This ROD describes disposal of contaminated materials at ERDF as the selected remedy, if the materials meet that facility's waste acceptance criteria (WAC). The purpose of this plan is to obtain characterization data that allow for proper disposal of the waste, whether at ERDF or another appropriately permitted facility. This document identifies the waste streams, as well as radiation survey and sampling approaches to be used to characterize the debris and soil. Throughout the text, this interim remedial action is referred to as a removal action or activity.

Figure 1-1 shows the location of the concrete pad and soil that will be removed. The personal protective equipment (PPE) is covered by other existing disposal documents and profiles and is not part of this SAP.



1.1 BACKGROUND

The K East Reactor and its associated fuel storage basin were constructed in the early 1950s. The basin is located in the Hanford 100 K Area within 1,380 ft of the Columbia River. The fuel basin is a large, open-topped concrete pool, containing approximately 1.3 million gallons of demineralized water. The basin was originally used to store SNF from the K East Reactor until the early 1970s, when the reactor was removed from service and the fuel removed from the basins. The K East basin subsequently was used to store SNF from the Hanford N Reactor. As of 1999, prior to removal of SNF from the basin, the K East fuel basins held approximately 1,200 metric tons of N Reactor SNF. The spent fuel elements are contained in canisters placed in storage racks under 16 ft of water for cooling and radiation shielding.

In the absence of a definition for debris in the 100 Area Remaining Sites ROD (EPA 1999), the definition given in the ERDF WAC (Bechtel 1998) will be used for this project. This definition is based on the *Resource Conservation and Recovery Act of 1976* (RCRA) definition of debris provided in 40 *Code of Federal Regulations* (CFR) 268.2 (g). For purposes of establishing disposal requirements, RCRA defines debris as a solid material exceeding a 60-mm particle size. The project does not anticipate that a significant quantity of material less than 60 mm will be generated. These items will generally be byproducts from and commingled with larger debris items and will be managed with the related waste stream.

Debris and soil management will depend upon the waste designation. All materials are anticipated to be low-level waste (LLW). This project is not expected to generate any transuranic (TRU) waste. Debris and soil might designate as hazardous or mixed waste, and may or may not contain polychlorinated biphenyls (PCBs).

The radiological historical data discussed in Section 1.1.1 apply to the concrete pad and soil. Filters containing residual water from the basin were previously stored on the pad, and thus may have contaminated the concrete surface with basin water. In addition, anecdotal descriptions indicate that basin water may have flowed to the soil outside the building before the pad was built.

The last paragraph of Section 1.1.1 explains the previous paint analyses that apply to the paint on the monorail and the painted portion of the concrete supports on the monorail. Note that the paint on the concrete filter wash pad is a new and different paint than that used on the monorail and associated monorail support concrete. The material safety data sheets (MSDSs) for the paint on the concrete filter pad are presented in Appendix B.

1.1.1 Previous Investigations

Summaries of historical data applicable to this project are provided below:

Mixed Waste Debris for the K Basins Project. An estimate of the Cs-137 content of the waste was performed for past shipments using established dose-to-curie relationships (WHC 1996a, WHC 1996b). Radionuclides considered reportable in previous waste shipments included strontium-90, Cs-137, plutonium-239/240, americium-241, and plutonium-241. The results of these Cs-137 estimates and the weights of the associated waste packages were used to develop a weight-to-curie relationship that will be applied to this waste (HNF 2001).

<u>Above – Water Waste</u>. Radiochemical analyses for gross alpha, gross beta, cobalt-60, Cs-137, and americium-241 were performed on twenty 105-KE smears. Nondestructive assay (NDA) of 20 compacted drums and NDA of four boxes of waste was performed. Radionuclides in the resulting waste profiles included strontium-90, Cs-137, europium-152, plutonium-238, plutonium-239, plutonium-240, americium-241, plutonium-241, and curium-244. The radiological data was used to develop an estimate of the radionuclide mix for above water waste in the K Basin (HNF 2000a, HNF 2001).

Fuel Basin Water. Because the basin water may have infiltrated the soil under and around the pad, and the residual water from the filters from the basin may have leaked on the concrete pad, the basin water characterization data are discussed here. Waters from the K East and K West Basins were analyzed for PCBs. PCBs were not detected in the water using a minimum detection limit of 0.5 ug/mL. Inductively coupled plasma (ICP) analysis for total metals was performed on water samples from both basins and on sludge from the K East Basin only. Although zinc, silicon, copper, and boron were detected in water samples, no toxicity characteristic (TC) metals were found above the TC levels, so the water is not a characteristic waste. Metals have been found in K East Basin sludge at concentrations that exceed the total concentration screening level. Only limited Toxicity Characteristic Leaching Procedure (TCLP) analyses were performed on the sludge. There are no data for mercury or organics. In addition, these existing TCLP data for the sludge cannot be correlated to the constituents in the water that may have accumulated in the soil.

Based on this information, the soil must be analyzed to assess whether characteristic metals are present. Because the water does not contain PCBs above the *Toxic Substances Control Act of 1976* (TSCA) levels of 50 ppm, and is not TSCA, the soil should not be a TSCA waste.

The above grade radionuclide ratios were developed in the previous SAP (HNF 2001). The source term for the above grade is most similar to the residual that may remain on the soil and the concrete pad. Note that the residual from air exposure around the basin is the same exposure method as the air around the monorail and monorail support. This is the reason that previous radionuclide ratios for above basin water are used in this SAP.

Paint Analysis applicable to monorail and concrete supporting the monorail. Analysis for total metals by ICP according to SW-846 Method 6010A [Test Methods for Evaluating Solid, Waste Physical/Chemical Methods, EPA 1997] were performed on nine paint chip samples, as well as multiple chip samples from an overhead crane. Toxic metals (silver, arsenic, barium, cadmium, chromium, lead, and selenium) were confirmed in paint chips at concentrations greater than screening limits for the TC criteria. The previous SAP (HNF 2001) discussed these data and the data was evaluated in a previous ERDF waste profile (#KBASIN001, Rev 01, 7/19/2001). The

TC constituent concentrations were divided by the total mass of the debris disposed, and the debris was not designated as characteristic waste in the previous waste profile. Therefore, this debris should not be designated as characteristic based on this information. The PCB data from the paint on the crane is presented in Appendix C and all concentrations are well below 50 ppm. Detection limits also are below 50 ppm. The SNF Project indicated the paint on the crane is of the same time frame as that used on the monorail and concrete supports. Based on the paint PCB data being below 50 ppm, this steel and concrete debris waste stream is not TSCA. The assessment is made on the paint alone and does not take into account the weight of the debris. Because a previous profile at ERDF exists for painted debris, the previous data supporting ERDF disposal (Profile #KBASIN001, Rev 01, 7/19/2000) will be used. No further analysis or evaluation of this waste stream is performed in this SAP.

1.1.2 Contaminants of Potential Concern

Constituents of potential concern (COPCs) include RCRA hazardous constituents (i.e., the TC metals and organics as listed in Table 2-2), applicable RCRA underlying hazardous constituents if the waste designates as characteristic, and PCBs as Aroclors. In addition, the ERDF WAC includes limits for additional organics and for radiological content.

The Data Quality Objectives (DQO) documentation for this project presents the rationale for exclusion of COPCs (EQM 2001). Table 1-6 in the DQO document (EQM 2001) provides the final list of constituents of concern (COCs) remaining for each waste stream with the rationale for inclusion and exclusion. The logic for selection of the radioisotopes is presented in HNF 2001 and EQM 2001. The radiological COCs are the same as those presented in the previous K Basin SAP (HNF 2001) for waste disposal at ERDF, and are listed in Table 2-3 of this document. The radiological COCs are based on the "above water" logic presented in HNF 2001 (see Appendix A). The logic is that water might have splashed and dried on the debris and soil, and thus represent the same source of radionuclides The soil underlying and surrounding the filter wash pad will be analyzed for the COCs listed in Table 2-3.

The debris waste streams (concrete and steel debris) will be evaluated based upon a previously employed weight-to-curie approach (HNF 2001) and will use the current ERDF Profile (#KBASIN001, Rev 01, 7/19/2001) for K-Basin debris.

For designation of the debris, the following approach has been used for the non-radiological COCs:

Concrete filter pad - Only a portion of the concrete pad is painted. The MSDSs for the paint are provided in Appendix B. The pad is relatively new, the paint is not the older paint previously used in the 100 Area, and the MSDSs do not list PCBs or metals as constituents. Therefore, PCBs are not a COC for the concrete. No characteristic metals or organics are listed on the MSDSs; therefore, no characteristic COCs apply.

<u>Soil</u> - No data exist for evaluation of the hazardous constituents in this soil; therefore, the soil will be analyzed for the TC constituent list and the additional organics required by the ERDF WAC.

Monorail and painted concrete supporting the monorail - The paint used for these areas was the same paint previously evaluated (HNF 2001). The data (see Appendix C) used for HNF 2001 for the paint indicated that it contained PCBs below the 50 ppm TSCA levels. The previous metals and organics data supporting the previous profile for debris from K Basin will be used to generate this profile.

<u>Transite siding</u> – This asbestos siding is to be removed from an external wall. It may or may not have been painted. If paint occurs, the time period to which the paint belongs has not been identified. Therefore, the waste stream will be evaluated for both old and new paints, based upon the previous evaluation for old paint (HNF 2001) and the MSDSs in Appendix B.

The asphalt is excluded by Washington Administrative Code 173-303-071 (3)(e) from analysis for disposal; therefore, no analyses will be requested for the asphalt and it is not given further consideration.

1.2 DATA QUALITY OBJECTIVES

The DQO process was employed to develop this SAP and determine the approach for characterizing the waste streams for disposal.

The scope of the DQO process (EQM 2001) included characterization of the five waste streams anticipated to result from this removal activity: (1) concrete pad debris and associated new paint; (2) underlying and surrounding soil; (3) asphalt from a road; (4) painted carbon steel monorail and concrete supports; (5) Transite siding from an external wall. The resulting information and data will allow the SNF Project to designate the waste streams for disposal. The DQO process was structured to provide the strategy for characterizing these waste streams in support of designation to determine the appropriate disposal facility and/or treatment required.

As noted above, decisions documented through the DQO process may be modified due to subsequent changes in project direction or based on discussions documented through the comment/response process. Any changes will be documented in project files. Refer to the DQO documentation (EQM 2001) for details about the process or resulting DQO.

1.2.1 Step 1: State the Problem

In the absence of a definition for debris in the 100 Area Remaining Sites ROD (EPA 1999), the definition given in the ERDF WAC (Bechtel 1998) will be used for this project. This definition is based on the RCRA definition of debris provided in 40 CFR 268.2 (g). Debris generated by remediation activities must be characterized and designated to allow disposal at ERDF, or be segregated for an alternate disposal pathway, as appropriate. All materials removed from this project area are assumed to be radioactively contaminated. Most debris will designate as LLW, although some may designate as mixed waste (i.e., radioactive and hazardous). For the radiological constituents in the concrete pad, the curie-to-weight method used in SAP (HNF 2001) will be used for the characterization for disposal.

Because no current data exist for the soil, soil underlying and outside the concrete structures will be excavated and data must be obtained to support designation for disposal.

1.2.2 Step 2: Identify the Decision

Step 2 presents the logic pathway that is used to resolve the problem. Tables 2-1 through 2-3 in the DQO document (EQM 2001) presents the Principal Study Questions, Alternative Actions, and Decision Statements to resolve the problem that was presented above. Figures 1-2 and 1-3 in this SAP present the decision logic, based on Step 2, which will be used to assess if the waste may be disposed at ERDF or other permitted disposal facility. This logic is the same logic previously used for waste designation for HNF 2001.

1.2.3 Step 3: Identify Inputs to the Decisions

Step 3 identified the data needed to resolve each of the Decision Statements identified in Step 2, as well as the analytical performance requirements (e.g., practical quantitation limits, precision, and accuracy) to support the data. The logic behind the selection of inputs, field techniques, and analytical methods and the tables, which present these information needs, may be found in the DQO document (EQM 2001). Because the MSDSs for the paint will be used to designate debris waste streams for TC metals and PCBs, no sampling and analysis will be conducted to support decisions related to these COCs. However, sampling and analysis will be performed to provide data to characterize the soil waste stream in support of waste designation. Table 2-2 presents a list of the radiological COCs and the radionuclide ratios that will be used to estimate the radionuclide content of the waste. Table 2-3 presents the list of radiological and non-radiological COCs and associated analyses to be used for the soil. Appendix A presents the final list of radionuclides to be addressed for the debris and soil waste streams. Appendix B presents the MSDSs for designation of the painted concrete pad.

Table 1-1 lists the decision statement, the waste stream and the associated data needed. For each set of decision, waste stream, and information needs, one or more existing documents are listed that allow a designation decision to be made. For soil, there is no data and, therefore, sampling is required. The table also provides general rationale and information with respect to waste characterization to support designation.

Table 1-1. Evaluation of Existing Data for Waste Designation (4 pages)

DS#	Waste Stream	Information :	Available Data	Is	Rationale				
	e graposti na producio di	Needed		information					
	- German Alberta (Company)	delegisie i Sprince e com		sufficient?	Congression of the second				
1,2, 3	Monorail,	Radiological	Above water	Y	Radionuclides source terms				
	concrete	composition	radionuclide		are expected to be the same as				
	support to rail,	(Curie/weight)	ratios from		K East above water debris				
	concrete pad,	and radionuclide	HNF 2001		1				
	Transite siding	ratios			•				
1,2,3	soil	Radiological	radionuclide	Ratios – Y	Radionuclides source terms				
		composition,	ratios from	Data to	are expected to be the same as				
		radionuclide	above water	confirm ratios	K East above water debris.				
		ratios '	HNF 2001, no	- N	Analyses will be performed				
			data available to		for gamma emitters and, if				

Table 1-1. Evaluation of Existing Data for Waste Designation (4 pages)

nous:			Existing Data for	Is S	Rationale
DS#	Waste Stream Fig. 1997 Bush Bush Street	Information Needed	Available Data	information sufficient?	Rationale
tinka: iks/ise (.a			confirm ratios		necessary, then for other radionuclides.
4	Monorail, concrete support to rail, concrete pad, Transite siding	Listed dangerous waste	No documentation of listed waste sources.	Y	No evidence exists to believe there were listed wastes (F, K, P, or U) used in association with the waste streams. EPA policy is to apply listed waste codes only when process knowledge indicates a listed waste source.
4	Monorail, concrete support to rail, concrete pad, Transite siding	Characteristic waste	NHC-96-101, "Analytical Report for K Basin Paint — FT-6112," 9/11/96. Analytical report for FAST Project FD1- 7021, "K Basin Crane Removal," 8/5/97. ERDF Profile #KBASIN001, Rev. 01, 7/19/01. Process knowledge.	Y	These materials are solids, which are not ignitable, do not support a corrosive solid (WA state) designation, are not explosive, and do not contain reactive sulfides or cyanides. Seven toxic metals (silver, arsenic, barium, cadmium, chromium, lead, and selenium) were confirmed to be present in paint chips at concentrations greater than screening limits for TC criteria. However, it is assumed that the paint remaining on the surfaces is less than 0.05% of the waste streams; therefore, the waste streams will not designate for toxic metals. Because pesticides/ herbicides are not in the paint formulations and the paint is dry (i.e., no volatile organics are present), none of the remaining characteristic codes will apply to these waste streams. The TC volatile organics were previously removed based on: Paint was applied over 30 years ago and, due to low vapor pressure, volatiles organics are no longer present. Data for paint in Appendix C shows no total halogens. As many volatiles are halogenated this provides added support that no volatiles
4	Monorail,	Toxic dangerous	ERDF Profile	Y	are present. Toxic waste constituents are

Table 1-1. Evaluation of Existing Data for Waste Designation (4 pages)

	Table 1-1. Evaluation of Existing Data for Waste Designation (4 pages)							
DS#	Waste Stream	Information Needed	Available Data	Is information sufficient?	Rationale			
	concrete support to rail, concrete pad, Transite siding	waste	#KBASIN001, Rev. 01, 7/19/01. Process knowledge	- 10 - 10 - 10 - 10 - 10 - 10 - 10 - 10	not suspected in these waste streams of steel, concrete, and asbestos materials. However, if toxic constituents are present in the paint and it is assumed that the paint remaining on the surfaces is less than 0.05% of the waste streams, the waste streams will not designate as toxic dangerous waste.			
4	Monorail, concrete support to rail, concrete pad, Transite siding	Persistent dangerous waste	ERDF Profile #KBASIN001, Rev. 01, 7/19/01. Analytical report for FAST Project FD1- 7021, "K Basin Crane Removal," 8/5/97. Process knowledge	Y	Halogenated organic compounds and polycyclic aromatic hydrocarbons are not suspected in the waste streams at concentrations that would cause the waste to designate as persistent dangerous waste.			
4	Painted monorail, concrete support to rail, Transite siding (old paint)	PCB concentrations	Analytical report for FAST Project FD1- 7021, "K Basin Crane Removal," 8/5/97. Process knowledge	Y	PCBs are not a constituent of the steel, concrete, or Transite, although some of the surfaces retain paint; this old paint is expected to be the same as found on the crane. PCBs have been found, associated with old paint, elsewhere on the Hanford Site at levels above 50 ppm but below 500 ppm.			
4	Concrete pad with (new) paint	PCB concentrations	MSDSs; see Appendix B	Y	MSDSs for new paint used in the 100 K East Area do not identify PCBs as an ingredient.			
4	Painted monorail, concrete support to rail, concrete pad, soil	Asbestos	Process knowledge	Y	These materials are solids which do not contain asbestos.			
4	Transite siding	Asbestos	Process knowledge	Y	This building material is known to contain asbestos.			
5	Monorail, concrete support to rail, concrete pad,	Land disposal restricted waste	Process knowledge	Y	Because the waste streams are not expected to designate, federal LDRs are not applicable. Also, the material			

Table 1-1. Evaluation of Existing Data for Waste Designation (4 pages)

DS#*	Waste Stream	Information	Available Data	Is	Rationale
		Needed		information sufficient?	
	Transite siding				is not liquid, a solid acid waste, or organic/carbonaceous and is not believed to be extremely hazardous waste; therefore the state LDRs do not apply.
4	Soil	Listed dangerous waste	No documentation of listed waste sources.	Y	No evidence exists to believe there were listed wastes (F, K, P, or U) used/generated in the 100 K Area. EPA policy is to apply listed waste codes only when process knowledge indicates a listed waste source.
4,5	Soil	Characteristic waste, toxic dangerous waste, LDR waste, PCBs	no data	N	Analyses will be performed. Designation will be based on the analyses.
4	Soil	Persistent dangerous waste	no data	N	Designation will be performed based on data from analyses by Methods 8260 and 8270.

The U.S. Environmental Protection Agency (EPA) policy regarding the application of a listed waste designation is that the listed waste codes should be applied only when the source of the waste is known and it can be concluded to require the application of the listed code. Even when analysis indicates the presence of a constituent that could carry a listed waste code, unless process knowledge indicates that the constituent was used for a listed purpose, the code should not be applied. There is no available information that indicates a listed waste code should be applied to the K Basin waste streams; therefore, no listed waste codes will be applied.

A radiological survey of the filter wash pad was performed on 7/28/00 (SNF Project Radiological Survey Report (K000511)). Although the data was not gathered to support waste characterization, it provides information as to the potential contamination levels that will be encountered during this project. HNF 2001a provides a calculation that supports that use of the weight to curie conversion factor currently used for K Basin debris.

1.2.4 Step 4: Define the Boundaries of the Study

Step 4 and Figure 1-1 identifies the geographic (spatial) and temporal boundaries of the area under investigation, as well as practical constraints that must be considered in the sampling design. No temporal boundaries have been identified. Table 4-1 in the DQO document (EQM 2001) defines the attributes that make up each population of interest. It is expected that approximately 40 cubic yards of concrete, 260 cubic yards of soil, 20 cubic yards of asphalt, and 270 square ft of Transite siding will be removed and require disposal from K East.

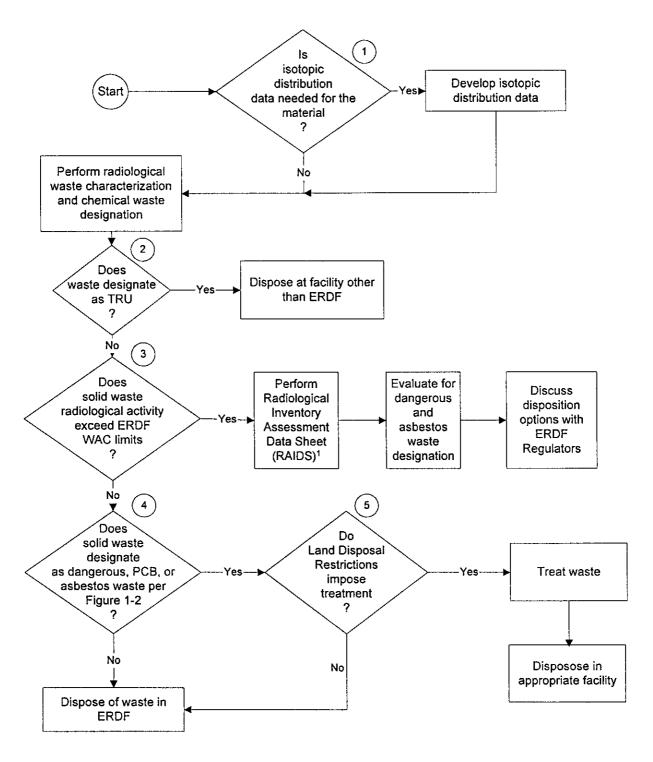
The geographic area of investigation includes the portion of the filter wash pad that is not currently marked as a radiation control area, and the underlying soil to a depth of at least 4.5 ft, external to the north side of the 100 K East Fuel Transfer System building. Decisions on the disposition of concrete pad or soil (i.e., the scale of the decision) will be made for the entire area being investigated.

The decisions identified in the DQO process apply to the removal of all debris or soil covered by this removal action. These decisions may or may not be appropriate for later debris or soil removal activities external to the K East buildings. The difficulty associated with collecting representative samples from debris waste streams supports use of the weight-to-curie conversion rather than sampling and laboratory-based analysis of radionuclides.

1.2.5 Step 5: Develop a Decision Rule

Step 5 combines information developed in DQO Steps 1 through 4 with a statistical parameter of interest (such as mean, median, or percentile) and an action level to provide a concise description of what action will be taken based on the results of the data collected. Table 5-2 in the DQO documentation (EQM 2001) lists the final action level for each Decision Statement and COC. This information is incorporated into the analytical performance requirements presented later in this SAP.

Figure 1-2. K East Area Debris and Soil Disposition Decision Logic



¹SNF staff will provide the necessary,inputs for the ERDF to perform calculations. It is not anticipated that the proposed waste will present any problems for the ERDF inventory.

Does Determine if solid waste contain known or solid waste exhibits toxicity suspect TC constituents characteristics Ν Apply process knowledge or waste a 100% solid perform TCLP or totals analysis on solids media Apply process knowledge or waste a liquid with < 0.5% perform TCLP or totals dry solid analysis on liquids Apply process knowledge or perform TCLP or totals Waste is a mixture of liquids and solids analysis using a formula for liquid/solid mixture Does waste Apply toxicity exhibit toxicity characteristic code characteristics Determine if designation per Washington Administrative Determine if solid waste is Code 173-303-100, Section 5 regulated for asbestos and/or 6, is required Return to Figure 1-1 decision diamond #4 1. Summarize codes for solid "yes" branch waste waste codes 2. If any codes apply, determine apply to LDR requirements, including solid waste application of UHC standards, Return to Figure 1-1 as appropriate decision diamond #4 "no" branch N

Figure 1-3. Chemical Waste Designation Decision Logic

Table 1-2 below (Table 5-3 in the DQO document (EQM 2001)) combines the parameter of interest, scale for decision making, action levels, and alternative actions into separate "IF...THEN..." Decision Rules. These decision rules are the output from the DQO process and describe actions that will be taken based upon the results of data analysis.

Table 1-2. Decision Rules

DS#	Decision Rule (2 pages)
1, 2, 3	If the radionuclide COCs in the waste do not exceed the radionuclide ERDF WAC (BHI 1998) (Ci/m³), then the waste will be evaluated by Decision Rules # 4a, 4b, and 5.
	If the radionuclide COCs in the waste exceeds the radionuclide ERDF WAC (BHI 1998) (Ci/m³), then the waste will be evaluated on a case-by-case basis for possible ERDF disposal, but may not be sent to ERDF. The waste will be evaluated by Decision Rules # 4a, 4b, and 5.
4a	If process knowledge or concentrations of the detected analytes indicate that the waste does not designate as listed, TC, toxic dangerous (i.e., state dangerous waste (DW), persistent dangerous (i.e., state dangerous extremely hazardous waste (EHW), PCB, or asbestos, then the waste will be evaluated for disposal at ERDF or another permitted facility.
	If process knowledge or concentrations of the detected analytes indicate that the waste designates as listed, TC, toxic dangerous (i.e., state DW, persistent dangerous (i.e., state dangerous EHW, PCB, or asbestos, then the waste will be evaluated for disposal at ERDF or another permitted facility.
4b	If process knowledge or concentrations of the detected analytes do not exceed the ERDF WAC limits, the waste will be disposed at ERDF.
	If process knowledge or concentrations of the detected analytes do exceed the ERDF WAC limits, then discuss with regulators the options of treating the waste and/or sending the waste to another permitted facility
5	If process knowledge or any detected analyte dictates LDR-required treatment and disposal, the waste will be treated and disposed at ERDF or another permitted facility.
	If no process knowledge or none of the detected analytes dictate LDR-required treatment and disposal, the waste will be disposed in ERDF or another permitted facility without treatment.

^{*}Radionuclide content is estimated from weight to Cs-137 curie conversions for debris, followed by application of the radionuclide ratios in Table 2-2. For soil, the Cs-137 and other gamma emitters will be obtained from laboratory data, followed by application of the radionuclide ratios in Table 2-2.

DS = decision statement

1.2.6 Step 6: Specify Limits on Decision Error

This section of a DQO generally is used to establish the parameters for a statistically-based sampling design. This SAP does not describe a statistically-based sampling approach for debris. Debris will be evaluated through weighing of all materials followed by use of the curie/weight and application of the above water radionuclide ratios in Table 2-2 as discussed in HNF 2001.

Soil will be evaluated for non-radiological COCs by random sampling designed to establish a mean estimated concentration for all soil to be removed. Radiological COCs for the soil will be estimated using a judgmental sampling of areas of maximum dose rate. Refer to Step 6 in the DQO document (EQM 2001) for additional details.

Contingency sampling will be performed only if anomalous waste is generated. The contingency sampling is discussed in Section 2.4.

1.2.6.1 Radioactive Debris Waste

Each waste container of debris will be weighed. An estimated COC inventory for waste containers of debris will be derived from weight-to-Cs-137 curie data coupled with radionuclide ratios from the previous K Basin SAP (HNF 2001).

1.2.6.2 Painted Debris

Designation for characteristic waste of the painted waste streams (concrete and metal debris from the monorail and concrete pad) will be based on the contribution of the hazardous constituents in the paint layer to the mass of debris being disposed.

For painted monorail and concrete supports, the PCB concentration in the paint will be used to designate with respect to PCB concentration.

1.2.6.3 Soil

Sampling and analysis of the soil underlying and adjacent to the concrete is planned to characterize the waste stream for designation. There are three options discussed for the timing of soil sampling: (1) sample before digging, (2) sample during the soil removal, and (3) sample from the bags after removal. The timing of sampling is important because the project must remove soil before fixed laboratory data will be completed and thus before a profile and ERDF roll-off (or other approved facility transportation boxes) will be obtained. The project will store soil in large plastic disposal bags until the soil is disposed at ERDF or other approved facility.

Option 1. This option will be used if samples are collected before digging. Triangular grids will be placed on the soil surface after removal of the pad. The grids will cover the soil under the pad and area adjacent to the pad that will be removed. For radionuclides, the surface soil will be surveyed, the three highest dose rate areas located, and subsequent samples in those three areas collected down to depth of excavation predicted for that area (4.5 ft to a few inches). Each sample will be a composite from the surface to a maximum depth of 4.5 ft. The depth is to be equivalent to the planned depth of excavation. This is a biased sampling design and will result in a conservative estimate of radionuclide concentration.

For the non-radiological parameters, there is no demonstrated association between (1) metals and organics and (2) the radionuclides. Therefore, three samples from randomly selected grid nodes will be collected. Each sample will be collected and composited from the surface to a maximum depth of 4.5 ft at each location. The depth of sampling is to be equivalent to the planned depth of excavation. The samplers will discuss with the construction engineers the depth of planned excavation before sampling.

Analysis for all options. For all options for rad analysis of soil, the following will be done. Initially, gamma energy analysis (GEA) of the samples for gamma emitters will be performed by the fixed laboratory. The ratios of individual gamma emitters to Cs-137 will be compared to the values in Table 2-2. If the ratios are less than or equal to those in Table 2-2, no additional analyses will be performed, and the ratios based on Table 2-2 will be used to estimate radionuclides other than Cs-137. If the gamma analysis indicates that the ratios are not consistent with the values in Table 2-2, then radionuclides in Table 2-3 may be analyzed and the actual concentrations used for all measured isotopes.

For all options for analysis of non-rad, the analytes, methods and practical quantitation limits (PQLs) in Table 2-3 will be obtained.

Option 2: This option will be used if sampling is performed during soil removal. In this case, it is assumed that the bulldozer or backhoe will be used both for excavation of the soil and for digging in areas to allow sample collection. Triangular grids will be placed on the soil surface after removal of the pad. The grids will cover the soil under the pad and area adjacent to the pad that will be removed. For radionuclides, the surface soil will be surveyed using the same grids previously discussed. The three highest dose rate areas will be located and marked with flags. As excavation proceeds, samples in those three areas will be collected down to depth of excavation predicted for that area (4.5 ft to a few inches). Each sample will be a composite from the surface to a maximum depth of 4.5 ft. The depth is to be equivalent to the planned depth of excavation. This is a biased sampling design and will result in a conservative estimate of radionuclide concentration. The radionuclide and non-radionuclide analyses will be the same as previously discussed.

For the non-radiological parameters, the same grids will be used. Three random locations will be selected and flagged before digging begins. Three samples from randomly selected grid nodes will be collected as the excavation proceeds. Each sample will be a composite from the surface to a maximum depth of 4.5 ft at each location. The depth of sampling is to be equivalent to the planned depth of excavation. The samplers will discuss with the construction engineers the depth of planned excavation before sampling.

Option 3. This option will be used if the samplers cannot be present during digging and the soil must be bagged before sampling. Triangular grids will be placed on the soil surface after removal of the pad. The grids will cover the soil under the pad and area outside the pad that will be removed. For radionuclides, the surface soil will be surveyed using same grids previously discussed. The three highest dose rate areas will be located and marked with flags. As excavation proceeds, for radionuclides, the soil from the three highest surface survey areas will be placed in one or more bags and will be marked for radionuclide sampling. A sample from each of three bags containing the high survey data surface soil will be obtained. Each bag contains about 9.5 cu yd and has a large opening to allow for obtaining the samples.

Workers removing the soil will keep a map and log identifying the location and depth from which each bag of soil is collected. Each bag will be randomly numbered; three random numbers will be selected for sampling the non-radionuclide analytes. A clean, small shovel or trowel or small hand coring device will be used to get samples from various soil portions within each bag. A sample will be collected from each of the three randomly numbered bags chosen for sampling.

The following tables show, under a variety of assumptions of what the sample standard deviation might be, what sample sizes would achieve the desirable error rates ($\alpha = 0.10$, $\beta = 0.20$). For each table, the regulatory limit of an inorganic analyte is taken as the Action Limit. The sample standard deviation(s) is varied to show the impact on sample size of the variability in the concentrations.

No analytical data exist for the concentration of any parameter to be measured in the soil underlying or adjacent to the concrete pad. Of the eight TC metals, arsenic, chromium, lead and silver (As, Cr, Pb, and Ag) have same toxicity characteristic action limit of 100 ppm (based on total analysis) and all have the same published variance; therefore, lead represents these metals. Lead is the logical choice because there are at least two possible sources of contamination. Vehicles which used leaded gasoline have driven on the soil, and lead is a known contaminant from vehicle exhaust. In addition, lead shielding was used in this area and is known to shed lead particles. Barium (Ba) has one of the higher limits and was selected for this reason. This leaves mercury, selenium, and cadmium (Hg, Se, and Cd) which have various action limits. There is no reason, given the volatility of mercury and selenium and the mobility of mercury, to suspect that these would remain in the soil. In addition, cadmium and the organic compounds are not expected to be present. Because it is more likely that lead and barium may be present in the soil, these metals were used to estimate the number of samples.

The following discussion shows, under a variety of assumptions of what the sample standard deviation might be, what sample sizes would achieve the desirable error rates ($\alpha = 0.10$, $\beta = 0.20$). In each table, the regulatory limit of an inorganic analyte is taken as the Action Limit. The sample standard deviation is varied to show the impact on sample size of the variability in the concentration.

For lead:

Action Limit = 100 mg/kg for the toxicity characteristic based on total analysis Δ = width of gray region (area of greater uncertainty and inability to determine with sufficient confidence that the site concentration is truly below the Action Limit)

Sample Size Determination for Lead Under Varying Assumptions

				****	<u> </u>
$(\alpha = 0.10, \beta = 0.20)$	s = 5	s = 10	s=20	s=30	
$\Delta = 20$	2	6	11		s = 13.9*
(20% of Action Limit)					3
$\Delta = 10$	2	6	19	42	s = 6.9*
(10% of Action Limit)					3

^{*}Greatest standard deviation for which three samples meet the desired error rates.

For barium:

Action Limit = 2000 mg/kg for the toxicity characteristic based on total analysis Δ = width of gray region (area of greater uncertainty and inability to determine with sufficient confidence that the site concentration is truly below the Action Limit)

Sample Size Determination for Barium Under Varying Assumptions

$(\alpha = 0.10, \beta = 0.20)$	s = 50	s = 100	s = 150	s = 300	
$\Delta = 400$	2	2	2	4	s = 278*
(20% of Action Limit)	1	ļ			3
$\Delta = 200$	2	2	4	11	s = 139*
(10% of Action Limit)	Į				3

^{*}Greatest standard deviation for which three samples meet the desired error rates.

The sample sizes under these two examples imply that, if we are comfortable with $\alpha=0.10$ and $\beta=0.20$ and with a gray region equal to 20% of the Action Limit, then three samples are sufficient to meet the error tolerances under assumptions that the standard deviation for lead will be 13.9 mg/kg or less, and that for barium will be no larger than 278 mg/kg. These standard deviations are greater than the expected analytical variation based on current precision estimates. The added variation allows for heterogeneity of the soil. These seem like reasonable assumptions for the expected concentrations, implying that three samples is a reasonable sample size to propose. In order to achieve low standard deviations in analytical data, it is recommended that sample handling and preparation carefully follow approved standard operating procedures, and that blending of the composited segment of the core sample be done as thoroughly as possible. (However, note that the samples taken for analysis of volatile organics will not be blended or homogenized. Blending or homogenization would cause loss of volatile analytes.)

1.2.6.4 Transite Siding

Transite siding is know to be made of asbestos and may have both old and new paint. It will be evaluated by process knowledge.

2.0 QUALITY ASSURANCE PROJECT PLAN

2.1 PROJECT MANAGEMENT

The following section identifies the individuals or organizations participating in the project and discusses specific roles and responsibilities. This section also discusses quality objectives for measurement data and discusses special training requirements for staff performing the work.

2.1.1 Project and Task Organization

Figure 2-1 presents the organization chart for sampling/analysis and waste management interfaces to ERDF.

2.1.2 Roles and Responsibilities

This section identifies the responsibilities of various organizations supporting this K East removal activity which collect, analyze, survey, or assess results of data for waste disposal.

K Basin Operations Support Sample Management Representative (contingency only)

- Used for contingency sample management only
- Maintain operating procedures as custodian, and revise procedures (if necessary) to perform sampling that include collection, chain of custody, packaging, and shipping procedure
- Maintain sample analysis records in a 2-hour-rated fire resistant file cabinet
- Receive data packages
- Perform or contract data review
- Maintain copies of radiological survey records and assemble into files to support waste characterization and designation

Fluor Hanford, Inc. (FH) Nuclear Process Operators (Duties apply to Contingency Sampling only; do not apply to soil.)

- Perform sampling for contingency sampling only
- Document sampling activities in a controlled logbook
- Initiate chain-of-custody documentation
- Package and ship samples to the specified laboratory

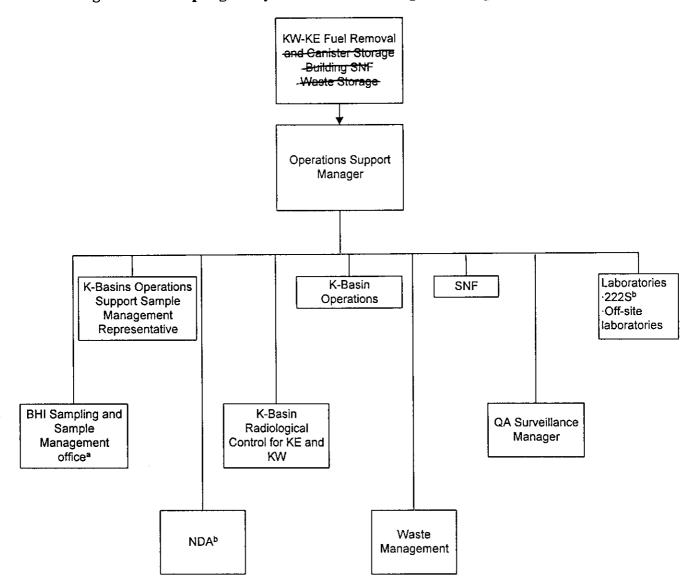


Figure 2-1. Sampling/Analysis and Waste Management Organization Chart

a BHI will collect, ship, and manage sample analysis other than contingency analysis

b Used for contingency analysis only

Specified Analytical Laboratory

- Adhere to and use procedures that are equivalent to those listed in Table 2-3, Sections 2.3, 2.6, 3.8, and 3.9
- Adhere to the latest revision of HASQARD
- Receive samples and initiate internal chain-of-custody documentation
- Provide specified radiological or non-radiological analyses
- Provide specified data package to the Operations Support Sample Management Representative

K Basin Operations Support Manager (or designee) (contingency samples only)

- Oversees sample management program
- Authorizes new radionuclide ratios to be applied to waste, as needed
- · Obtain additional analytical services such as NDA, if needed

FH Radiological Control Organization

- Conduct specified surveys and/or NDA
- Provide dose rate data for sample collection, packaging, and transportation or shipment to the laboratory used for contingency analysis discussed in Section 2.4
- Provide the Radiological Work Permit

Waste Management FH

- Review characterization data used to designate waste
- Provide shipping documentation to ERDF

ERDF Waste Management (Bechtel Hanford, Inc.[BHI])

- Designate waste based on survey/laboratory results and calculated radionuclide content
- Supply shipping containers for waste going to ERDF
- Dispose of waste that meets ERDF WAC in ERDF

Quality Assurance Organization

• FH quality assurance (QA) and BHI QA have the option to conduct random surveillances to verify compliance with requirements of this plan

Environmental Restoration Contractor (ERC) Analytical and Field Services

- CH2M Hill Hanford/BHI will collect only the soil samples
- Manage the logistics of obtaining soil analyses from offsite laboratories
- BHI will not perform contingency sampling or analyses

2.1.3 Special Training Requirements/Certification

Hazards associated with radiation and radiological contamination are well known and documented for the K East Area. The *Hanford Site Radiological Control Manual* (HSRCM) (DOE-RL 1996) addresses worker training requirements, visitor training and escort requirements, dosimetry, monitoring, posting, and required radiological surveillance. The specific training required by 29 CFR 1910.120 is implemented by the HSRCM. Training requirements for this project are discussed in Section 7 of the *K-Basins Interim Remedial Action Health and Safety Plan* (HASP) (HNF 2000b). Project-specific training requirements and references are discussed below.

In the event that a worker may have a reasonable possibility of exposure to hazardous chemicals while performing a specific remediation task in the K East Area, the worker's supervisor will ensure that the worker has the appropriate level of training, in accordance with 29 CFR 1910.120, before the work is performed.

SNF Project Administrative Procedure AD-14-004, "Radiological Area Access Control," defines training requirements for various circumstances applicable to entry into K Basins. Training requirements in this procedure apply to all individuals who are required to have access to the K Basins.

Job-specific training requirements for SNF Project personnel are outlined in Procedure TN 8-001-08, "General Training Administration." This procedure covers facility orientation training, Hanford General Employee Training, facility emergency plan, SNF Project orientation, initial and continuing training, on-the-job training, required reading and drills. The training requirements for each employee are determined using a graded-approach and are documented in the appropriate Training Matrix.

All visitors, general employees, or members of the public, will have training or instruction before entry to the K Basins according to the requirements of SNF AD-14-004.

2.1.4 Quality Objectives and Criteria for Measurement Data

The QA objective of this plan is to develop implementation requirements that will provide data of known and appropriate quality. Data quality is typically assessed by representativeness, comparability, accuracy, precision, and completeness. These parameters are described below. The applicable quality control (QC), quantitative target limits, and levels of effort for assessing data quality are dictated by the intended use of the data and the nature of the analytical method. A summary of COCs for each waste stream is provided in Table 1-6 of the DQO document (EQM 2001). The analytical methods, laboratory detection limits, and test portion for COCs that will be measured are presented in Table 2-3 for soil samples. Table 2-4 provides the same information for contingency samples. The COCs that are not listed in these tables will be estimated based on radionuclide ratios in the waste as discussed in Section 2.2. QC parameters of accuracy and precision that will be applied to soil or contingency characterization samples are presented in Table 2-4. The nomenclature used to describe quality parameters is discussed below.

Representativeness is a measure of how closely measured results reflect the concentration of constituents distributed in the sample matrix. Sampling plan design, sampling techniques, and sample handling protocols (e.g., storage, preservation, and transportation/shipping) have been developed and are discussed in later sections of this document. Documentation will establish that protocols have been followed and sample identification and integrity are ensured.

Comparability expresses the confidence with which one data set can be compared to or used with another data set. Data comparability will be maintained by using standard documented procedures, consistent methods, and data reporting units. Analytical methods and target detection limits are listed in Table 2-3. Actual detection limits will depend on the sample matrix, constituent, and test portion and will be reported as defined for the specific samples. Detection limits are functions of the analytical method utilized to provide the data and the test portion analyzed. For the soil analyses and contingency sampling, sufficient amounts of sample material are expected to be available. In addition, for radiological analyses, sufficient radionuclide activity to perform the analyses also is expected.

Accuracy is an assessment of the closeness of the measured value to the true value. Accuracy of chemical test results is assessed by spiking samples with known standards and establishing the average recovery. A matrix spike is the addition to a sample of known amounts of a standard compound similar to the compounds being measured. Radionuclide measurements that require chemical separations often use this technique to measure method performance. For radionuclide measurements that are analyzed by gamma spectroscopy, laboratories typically compare results of Laboratory Control Samples against known standards to establish accuracy. Usually, only a few target analytes are selected for QC analysis for gamma spectroscopy (e.g., Cs-137, cobalt-60). Calibrations are evaluated by comparing results from measurements of check standards to known values and/or by generation of in-house statistical limits. Table 2-3 lists the accuracy targets for laboratory analyses for the project.

Precision is a measure of the data spread when more than one measurement has been taken on the same sample. Precision can be expressed as the relative percent difference for duplicate measurements or as the relative standard deviation for other multiples. Precision targets for laboratory analyses are listed in Table 2-3.

Completeness is a comparison of the valid data required to the amount of valid data obtained from the analytical measurement process and the complete implementation of defined field procedures. The completeness objective for this SAP is 100%. Completeness will be assessed by waste stream on an analyte-specific basis. If the completeness objective is not met, additional samples will be collected and analyzed.

2.1.5 Documentation and Records

Field logbooks contain area and task-specific information. Field logbooks that are used during collection of samples for waste characterization will be identified as a quality record and will be maintained as such.

Maintenance of field documents will be in accordance with the *Hanford Analytical Quality Assurance Requirements Documents* (HASQARD) (DOE-RL 1998) Volume II, "Sampling Technical Requirements," Section 4.1.2, "Field Logbook," or equivalent.

2.2 SURVEY/DATA ACQUISITION

The following sections present the logic and requirements for radiological survey. The radiological dose rate survey data will be used to guide selection of sample points in the underlying soil after removal of the concrete pad.

Waste generated by this removal activity and deemed appropriate for shipment to ERDF will be processed to comply with ERDF WAC (BHI 1998) and packaged according to Procedure OP-46-006 ("Processing Contaminated Waste for ERDF Disposal").

Packaged waste will be surveyed according to appropriate instrument procedures to assure that the outside of the waste debris /soil package meets surface contamination limits. The survey will be documented per HNF-PRO-1892 ("Documentation of Radiological Surveys"), and shipped to ERDF where it will be weighed. The weight of the waste package will be used to estimate the Cs-137 curie content of the debris waste streams as discussed below. Soil will be analyzed for Cs-137 curie content as discussed in Section 2.3. Utilizing the ratios of the COCs to Cs-137 as discussed in Section 2.2.1, the radionuclide content of both debris and soil will be estimated.

If the waste is determined to be anomalous (as defined in Section 2.2.3), it will be set aside and may be subjected to sampling and analysis. The approach for contingency sampling is discussed in Section 2.4 of this SAP. The sections below address requirements for instrument calibration and maintenance, and data management.

2.2.1 Debris Weight to Cs-137 Curie Conversion

The weight-to-curie relationship that will be utilized for K Basin waste is based on the dose-tocurie relationships found in methods discussed in WHC-SD-WM-PROC-020 ("Procedure for Categorizing and Inventorying Waste in Standard Containers" [WHC 1996e]). The technical basis for this procedure is presented in WHC-SD-WM-RPT-267, "Basis for Dose Rate to Curie Assay Method" (WHC 1996a). Briefly, the method utilizes a family of curves that relate the measured dose rate (R/hr) to the Cs-137 curie content of the waste. Although the technical basis document (WHC 1996a) was prepared for tank waste, the basic premise of the document is that the major contributor to the measured dose rate is Cs-137. That same premise is appropriate for the K Basin debris. Although other gamma emitters do exist in the K Basin debris, the most common (cobalt-60, europium-152, europium-154, and europium-155) generally are less than 10% of the Cs-137 content. By using the conservative assumption that all measured dose rate is from Cs-137, other gamma-emitting radionuclides, if present, would be led to an overestimation of the Cs-137 content of the waste. All other radionuclides will be estimated based on use of specific ratios of COC radionuclides to Cs-137 for the waste in question. Thus, the final estimated radionuclide content would likely be overestimated if gamma-emitting radionuclides were present in greater abundance than anticipated.

A review of the past K Basin waste disposal records was performed to develop a correlation between Cs-137 concentrations and the weights of the various waste streams, SNF-7895, "Documentation of K Basin Waste Determination Based on Cs-137 Concentration in Ci/Kg" (SNF 2001). This relationship was then utilized with the scaling factors from the SAP to determine the radionuclide concentrations.

The review consisted of looking at waste data from 1996 through early 2000 and comparing Cs-137 concentrations from each of the waste streams from K East and K West. There was no information available for K West below water debris and the information for K East below water debris did not separate washed and non-washed materials.

In late 1999, a change in the nuclide determination methodology was made and the dose-to-curie models were updated. This resulted in three orders of magnitude increase in waste Cs-137 concentration. In reviewing the data, the higher Cs-137 information was used to make the comparison.

The basis for the Cs-137 concentrations applied to the waste determinations prior to mid-1999 came from the WHC-SD-NR-RPT-005, "Characterization of Radioactive Waste at 100-N," November 1990 (WHC 1990). Methodology after mid-1999 utilized the methodology identified in WHC-SD-WM-RPT-267, "Basis for Dose to Curie Assay Method," October 1996 (WHC 1996a). This later method is recommended in the draft, "Sampling and Analysis Plan for K-Basins' Debris" (HNF 2000e).

The waste containers chosen to make the Cs-137 Ci/kg relationship were those that the Cs-137 concentrations in the waste were developed after mid-1999.

The methodology used to develop a Cs-137 waste concentration was accomplished by performing the following:

Waste information was obtained from the Solid Waste Information Tracking System and placed on an Excel spreadsheet. The data was then sorted to just provide Cs-137 waste concentration. The data was then further sorted into three waste streams, K East Above Water, K East Below Water and K West Above Water.

The K East and K West Above Water Waste was then sorted into those containers classified before September 1999 and those classified after September 1999. This allowed a statistical review of the data.

The mean standard deviation for Cs-137 and container weight of each set of data was determined. A 95% confidence level of the data was used to identify an upper limit for the waste. The Cs-137 concentration was determined to be the mean Cs-137 concentration of the waste plus 1.96 standard deviation of the Cs-137 concentration. The Cs-137 concentrations are presented in Table 2-1.

The Cs-137 concentration was then used to determine the COCs radionuclides using scaling factors identified in the "Sampling and Analysis Plan for K-Basin's Debris" (HNF 2001). These radionuclides were then compared against the ERDF WAC.

Waste Stream	Cs-137 Concentration (Ci/Kg)
K East Above Water	1.12 E-5
K East Below Water Washed Metal	1,46 E-4
K East Below Water Unwashed Metal and Non-Metal	1.46 E-4
K West Above Water	8.51 E-6
K West Below Water	1.46 E-4

Table 2-1. Cs-137 Concentration in K Basin Waste Streams

2.2.2 Cs-137 Curie-to-Radionuclide Content Estimate for Waste

In previous work (HNF 2001), a final list of COCs was generated for waste disposal from K East and K West. The logic and approach for selecting the final list of COCs is discussed in Appendix A of the K East Removal DQO (HNF 2000e) and Appendix A of this document. The COCs appropriate for this removal action at K East would be the same as those estimated for the above water waste from the K East Basin. As discussed in the previous document (HNF 2000e, HNF 2001), the source of radionuclides in the above water waste in the K Basin is ultimately N-Reactor fuel rods. The mechanism for deposition of contamination on above water waste is also discussed and is mainly dried Fuel Basin water and incidental sludge particles that might adhere to equipment and filters that are removed from the basin.

As discussed in the DQO for this project, the source of radionuclides on, around, and under the pad is basically the same as for the K East above water debris and the same list of COCs will be used. For debris waste streams, the Cs-137 content will be estimated through weight-to-curie conversions that are discussed above. The ratios of various COCs to Cs-137 have been estimated

based on review of available analytical data and computer calculations of estimated content of fuel and sludge from the K East Basin (HNF 2001). Table 2-2 lists the calculated ratios. The same ratios are assumed to apply to this project. The following sections discuss the use of estimated ratios to characterize waste.

2.2.3 Anomalous Waste

Two anomalies may result. One is based on visual examination of the debris and the other is based on the concentrations radiological and non-radiological contaminants that result after applying the methodologies (e.g., weight-to-curie estimates) or analytical data results to the waste.

A visual examination of the debris will be performed. If anything other than concrete debris including rebar, steel debris, Transite siding debris, soil, or asphalt is included, it will be set aside and the survey and sampling approach will be revisited. An alteration in the waste profile may be needed.

The radiological and non-radiological data obtained on the soil will be evaluated. For radiological data, if the ratios of individual gamma emitters (Co-60, Sb-125,Eu-152, Eu-154, Eu-155, Am-241) to Cs-137 exceed the percentages listed in Table 2-2, the soil samples will be analyzed for the remaining radionuclides listed in Table 2-3. In addition, the waste may be considered anomalous. The percentages listed in Table 2-2 represent the highest percentage of ratio from available data that was applicable to the K East above water debris, thus all of the current waste should fall within that profile. The anomalous waste will be segregated until the additional radionuclide data are available. Alternative survey/sampling and profiling will be performed as needed.

It should be noted that professional judgment will be used to evaluate the GEA data with regard to detection limits and analytical error. If it is determined that the observed exceedance of ratios in Table 2-2 is likely due to random statistical fluctuations as a result of low levels of radionuclide in the waste, no additional analyses or revision of ratios will be performed.

Table 2-2. Summary List of Radionuclide Contaminants of Concern and Ratios to Cs-137 for K East Debris and Soil (2 pages)

Radionuclide Name	Radionuclide Symbol	Chosen Ratio for K East Debris and Soil
Tritium	H-3	0.090%
Cobalt	Co-60	1.0%
Nickel	Ni-63	0.34%
Strontium	Sr-90	103%
Antimony	Sb-125	0.16%
Cesium	Cs-137	100%
Promethium .	Pm-147	2.3%
Samarium	Sm-151	1.4%

Table 2-2. Summary List of Radionuclide Contaminants of Concern and Ratios to Cs-137 for K East Debris and Soil (2 pages)

Radionuclide Name	Radionuclide Symbol	Chosen Ratio for K East Debris and Soil
Europium	Eu-152	0.062%
Europium	Eu-154	1.4%
Europium	Eu-155	0.45%
Uranium	U-234	0.027%
Uranium	U-235	0.0046%
Uranium	U-238	0.021%
Plutonium	Pu-238	2.1%
Plutonium	Pu-239	13%
Plutonium	Pu-240	5.1%
Plutonium	Pu-241	197%
Americium	Am-241	17%
Curium	Cm-244	0.013%

In addition to the gamma ratios, any waste for which the estimated radionuclide content is greater than 100 nCi/g total TRU or greater than Class C limits (10 CFR 61.55) will be set aside for possible sampling and analysis. The methodologies for estimating the individual radionuclides (weight-to-curie and soil analysis) are considered sufficiently conservative such that the decision level will be 100 nCi/g total TRU radionuclides. If sampling is required, the path forward must be developed because this contingency is not within the scope of the DQO process for this K East removal project. The path forward will have input from EPA, DOE, FH, and ERDF representatives. If a more precise measurement of the waste is not obtained, or if measurements confirm that the waste is potentially TRU, the waste cannot be disposed at ERDF and an alternate disposition pathway must be identified.

2.2.4 Radiological Survey Methods/Quality Control Requirements

Surveys of the surface of the waste packages will be performed (as appropriate) to the criteria discussed in Procedure OP-46-006 to determine if waste packages can be removed from the initial staging area and placed in a bulk waste container. Radiological protection technicians perform surveys and obtain smears from the surfaces of waste packages (typically wrapped or bagged in plastic) to assess compliance with the criteria in Procedure OP-46-006. It is anticipated that due to contamination levels on the waste and the general background in the bagging area, smears of waste package surfaces may be required before removal from the staging area. Appropriate scan speeds, survey techniques, and smear counting procedures are referenced in the instrument specific procedures that will be used (see Section 2.2.5.1).

2.2.4.1 Radiological Surveys

Radiological surveys of the outside of waste packages for radiological control purposes and to comply with ERDF waste surface contamination acceptance criteria will be performed as appropriate and reported according to the following procedures:

- OP-46-006
- HNF-PRO-1892

The instruments used will be as required by the procedures:

- Beta/gamma survey meter, "GM Portable Survey Instrument," HNF-PRO-632
- Dose rate meter, "Eberline RO-3B (CP)," HNF-PRO-648
- Alpha survey meter, "Portable Alpha Meter," HNF-PRO-633

2.2.4.2 OC Requirements for Radiological Surveys

QA is necessarily built into each phase of the characterization as field instrument operational checks that monitor field instrumentation performance.

Alpha, beta/gamma surveys, and dose rate measurements may be used. Instruments will be calibrated against known standards representative of the instrument response to the identified analyte. The instrument will be within the calibration period specified by the instrument procedure.

QC measures taken to support field operations performance, including daily calibration checks, which will be performed and documented on each instrument used to survey or characterize waste. These checks will be performed as defined in the appropriate instrument procedure.

2.2.4.3 Instrument Testing, Inspection, and Maintenance Requirements

Measurement and testing equipment used in the field will be subject to acceptance testing and preventative maintenance measures to ensure minimization of measurement system downtime. Maintenance requirements, such as parts lists and instructions, and documentation of routine maintenance, will be performed according to the general program procedure ("Radiation Protection Instrument Program," HNF-PRO-436), as well as any additional measures that are specified in the specific instrument procedure referenced in Section 2.2.6.1.

2.2.4.4 Instrument Calibration and Frequency

Instruments used for surveys and screening for off-site sample shipment will be calibrated in accordance with HNF-PRO-436. The results from all instrument calibration activities shall be recorded as defined in the program procedure. Control documents must specify when the instrument was last calibrated, the results of that calibration, and the due date for new calibration.

2.2.4.5 Inspection/Acceptance Requirements for Supplies and Consumables

Procurement activities related to radiological survey will be limited to performing acceptance testing for all instruments and standards used as described in the program procedure HNF-PRO-436 and specific instrument procedures.

2.2.4.6 Field Survey Documentation

Field documentation will be kept in accordance with HNF-PRO-1892.f

2.2.5 Waste Handling and Custody Requirements

All waste handling, shipping, and custody requirements will be met in accordance with Procedure OP-46-006, "Processing Contaminated Waste for ERDF Disposal."

In addition, radioactive waste will be surveyed for shipment in accordance with HNF-PRO-1892. Radiological survey tags will be attached to individual bags of waste until they are placed in a larger container, such as a drum or box. The survey tags from bags that go into a larger container will be retained in order to provide a record of the surveys and associated estimate of curie content of the waste.

2.2.6 Waste and Sample Packaging for Shipping/Transportation

Waste and contingency sample packaging for shipping/transportation will be performed in accordance with procedure OP-46-006, "Processing Contaminated Waste for ERDF Disposal." The contingency samples or waste will be shipped/transported according to the procedure HNF-PRO-157, "Radioactive Material/Waste Shipments."

The following applies only to contingency samples discussed in Section 2.4. The current process is that for samples transported to onsite laboratories, unused samples are sent back to the generator. For samples shipped offsite, unused samples are not returned. The contracts with offsite laboratories specify that the laboratory disposes of any remaining sample and the waste associated with analysis.

Section 3.0 discusses BHI procedures for shipping samples that apply to soil.

2.3 K BASIN SOIL SAMPLING FOR WASTE DESIGNATION

2.3.1 Sampling Approach for Radiological Analyses

Soil sampling will occur after removal of the concrete pad. Since anecdotal information indicates that the pad was placed on soil that contained low-level radiological contamination, the approach is biased to sample the areas of highest radiological contamination. Initially a dose rate or other applicable survey of the surface soil will be performed. The survey will use the same grid that is established for non-radiological sampling. Survey results will be reported on a standard SNF Project Radiological Survey Report Form (BD-6002-600).

At each of three locations containing the highest instrument reading (e.g., mR/hr), a sample will be collected to be analyzed as discussed in Section 3.1. The details of sample collection are discussed in Section 3.0. Because soil from the surface to a maximum 4.5 ft depth will be removed for disposal, either a shovel, hand auger, backhoe or bulldozer will be used for sample collection through the entire interval for options 1 and 2, and a clean shovel used for each of three samples for option 3, followed by homogenizing the soil and bottling for shipment to the standard fixed laboratory (SFL). This approach will be repeated for each sampling location.

2.3.2 Sampling Approach for Non-radiological Analyses for Soil Samples

The details of sample collection are discussed in Section 3.0. For all three options, all non-rad analyses will be the same and will be those analytes, methods, and PQLs listed in Table 2-3.

For the samples for the non-radiological analytes (i.e., metals, volatile organic compounds (VOCs), semi-volatile organic compounds, pesticides, and herbicides), a triangular grid will be drawn and random samples collected (Options 1 and 2) or one sample will be collected from each of three randomly numbered bags (Option 3). Past data from soils in the 100 Area have indicated no correlation between radiological and non-radiological contaminants; therefore, a random sampling will be representative of the waste.

For samples for all analytes except VOCs, the following will be done at each sampling point. Because soil from the surface to a maximum 4.5 ft depth will be removed for disposal, either a shovel or hand auger will be used for sample collection, followed by homogenizing the soil from the surface to a depth of 4.5 ft or homogenizing subsamples from a given bag and bottling for shipment to the SFL.

For samples for VOCs, from each shovel or hand auger, small portions of soil from throughout the interval to be excavated (i.e., surface to 4.5 ft) or small portions from the bag will be placed in the jar for volatile organic analyses. No homogenizing will be performed.

2.3.3 Soil Sample Handling and Custody Requirements

Sample handling, shipping and chain-of-custody requirements are discussed in Section 3.0.

2.3.4 Soil Sample Preservation, Containers and Holding Times

Preservatives, containers, and holding times are discussed in Section 3.0.

2.3.5 Soil Sample Shipping

Requirements are described in Section 3.0.

2.3.6 Soil Survey Radionuclide Survey Measurements

Requirements are described in Section 3.4

2.3.7 Standard Fixed Laboratory Measurements for Soil Samples

The laboratory shall adhere to HASQARD and shall adhere to chapter 1 of SW-846 and the QC described in the SW-846 methods. Exceptions to the two cited documents shall be pre-approved by Fluor and by ERDF before analyses shall begin.

Parameters for soil analysis are listed in Table 2-3. Laboratory-specific SOPs for analytical methods are in place. Laboratory SOPs and QA Plans to be used include Analytical Procedures and QA Plans that have previously been reviewed and approved by BHI contracts with off-site analytical laboratories. Changes or additional methods identified during future engineering or planning will be presented in page changes, addenda, or revisions to this SAP as appropriate. Detection limits achievable by the laboratory will be dependent on sample quantity available and also may be affected by the matrix and radionuclide activity levels of the sample. Two of the radionuclide COCs (Pm-147 and Sm-151) are not listed in Table 2-3 because there are no established analytical procedures to analyze them. These radionuclide COCs will be estimated based on the established COC ratios to Cs-137.

For all organic laboratory analyses, actual standards of each compound or Aroclors will be used to calibrate the equipment. The number of concentrations of the standards used in the calibration curve will be that required by SW-846 method cited in Table 2-3. Tentatively Identified Compounds analysis cannot be substituted for the use of the actual calibration standards required by the organic method specified in Table 2-3.

Table 2-3. Analytical Performance Requirements (3 pages)

0.966			A PALES		hideling a fabili	Requiremen	ts
Waste Stream	tream Concern Action Level total analysis		Analytical Method ¹ / Technique	PQL	Accuracy as Spike Recovery (%)	Precision as RPD (%)	
		mg/Kg	mg/Kg		mg/Kg		
	PCBs ²	500	500	8082	0.0165	70 - 130	<u>+</u> 30
Soil	HOCs (halogenated organic compounds) ³	<1000	<1000	NA	NA	NA	NA
And	Radionuclides ⁴	pCi/g⁴			pCi/g		
Soil	H-3 (tritium)	NL	NA	LSC	400	70 - 130	<u>+</u> 30
	Co-60	NL	NA	GEA	0.05	NA	± 30
	Ni-63	4.38E+08	NA	Sep/LSC	30	70 - 130	± 30
	Sr-90	4.38E+09	NA	GPC	1	70 - 130	± 30
	Sb-125	NL	NA	GEA	0.3	70 - 130	± 30
	Cs-137	2.00E+07	NA	GEA	0.1	70 - 130	<u>±</u> 30
	Eu-152	1.31E+13	NA	GEA	0.1	NA	± 30
	Eu-154	NL	NA	GEA	0.1	NA	± 30
	Eu-155	NL	NA	GEA	0.1	NA	± 30
	U-234	4.64E+04	NA	AEA	1	70 - 130	<u>+</u> 30

Table 2-3. Analytical Performance Requirements (3 pages)

						Requirements		
Waste Stream	aste Constituent of Action Level L		Action Level for total analysis	Analytical Method ¹ /- Technique	PQL	Accuracy as Spike Recovery (%)	Precision as RPD (%)	
	U-235	1.69	E+03	NA	AEA	1	70 - 130	± 30
	U-238	7.501	E+03	NA	AEA	1	70 - 130	± 30
	Pu-238	1.00	E+05	NA	AEA	1	70 - 130	± 30
	Pu-239	1.81	E+04	NA	AEA	1	70 - 130	<u>+</u> 30
	Pu-240	1.81	E+04	NA	AEA	1	70 - 130	<u>+</u> 30
	Pu-241	3.88	E+06	NA	LSC	15	75 - 125	± 30
	Am-241	3.12	E+04	NA	AEA	1	70 - 130	± 30
	Cm-244	1.00	E +05	NA	AEA	1	70 - 130	± 30
	TC metals ⁵	TC mg/L	LDR TCLP	mg/Kg ^{5,6}		mg/Kg		
Mit (Constitute 1 1 1 1 1	As	5.0	5.0	100	6010	10	70 - 130	<u>+</u> 25
Soil	Ba	100.0	21.0	420	6010	20	70 - 130	± 25
	Cd	1.0	0.11	2.2	6010	0.5	70 - 130	± 25
	Cr	5.0	0.6	12	6010	1	70 - 130	± 25
	Pb	5.0	0.75	15	6010	10	70 - 130	± 25
	Hg	0.2	0.025	0.5	7471	0.2	70 - 130	<u>+</u> 25
	Se	1.0	5.7	20	6010	10	70 - 130	± 25
	Ag	5.0	0.14	2.8	6010	2	70 - 130	± 25
		TC	LDR					Sa Greek
	TC organics ⁵	mg/L TCLP	mg/Kg	mg/Kg ^{5,6}		mg/Kg		
Soil								
	Endrin	0.02	0.13	0.13	8081	0.0033	70 - 130	± 30
	Lindane	0.4	0.066	0.066	8081	0.00165	70 - 130	± 30
	Methoxychlor	10	0.18	0.18	8081	0.0165	70 - 130	<u>+</u> 30
	Toxaphene	0.5	2.6	2.6	8081	0.165	70 - 130	<u>±</u> 30
	2,4-D (2,4- Dichlorophenoxy- acetic acid)	10	10	10	8151	0.4	70 - 130	± 30
!	2,4,5-TP (Silvex)	1	7.9	7.9	8151	0.02	70 - 130	± 30
	Benzene	0.5	10	10	8260	0.005	70 - 130	± 30
	Carbon tetrachloride	0.5	6	6	8260	0.005	70 - 130	± 30
	Chlordane	0.03	0.26	0.26	8081	0.0165	70 - 130	± 30
	Chlorobenzene	100	6	6	8260	0.010	70 - 130	± 30
	Chloroform	6	6	6	8260	0.05	70 - 130	<u>+</u> 30
	o-Cresol	200	5.6	5.6	8260	0.33	70 - 130	± 30
	m-Cresol	200	5.6	5.6	8260	0.33	70 - 130	± 30
	p-Cresol	200	5.6	5.6	8260	0.33	70 - 130	± 30
	Cresol (total)	200	na	11.27	8260	NA	70 - 130	± 30

Table 2-3. Analytical Performance Requirements (3 pages)

	Hamilia i la						Requiremen	ts
Waste Stream		Analytical Method ¹ / Technique	PQL	Accuracy as Spike Recovery (%)	Precision as RPD (%)			
	1,4- Dichlorobenzene	7.5	6	6	8260	0.33	70 - 130	<u>±</u> 30
	1,2-Dichloroethane	0.5	6	6	8260	0.005	70 - 130	± 30
	1,1- Dichloroethylene 2,4-Dinitrotoluene	0.7	6	6 2.6	8260 8270	0.01	70 - 130 70 - 130	± 30 ± 30
:	Heptachlor (and epoxide)	0.008	0.066	0.066	8081	0.00165	70 - 130	± 30
	Hexachlorobenzene	0.13	10	2.6	8270	0.33	70 - 130	<u>+</u> 30
	Hexachloro-1,3- butadiene	0.5	5.6	5.6	8270	0.33	70 - 130	<u>+</u> 30
	Hexachloroethane	3	30	30	8270	0.33	70 - 130	± 30
	Methyl ethyl ketone	200	36	36	8260	0.01	70 - 130	± 30
	Nitrobenzene	2	14	14	8270	0.33	70 - 130	± 30
	Pentachlorophenol	100	7.4	7.4	8270	0.33	70 - 130	± 30
	Pyridine	5	16	16	8270	0.66	70 - 130	± 30
	Tetrachloroethylene	0.7	6	6	8260	0.005	70 - 130	± 30
	Trichloroethylene	0.5	6	6	8260	0.005	70 - 130	<u>+</u> 30
	2,4,5- Trichlorophenol	400	7.4	7.4	8270	0.33	70 - 130	± 30
	2,4,6- Trichlorophenol	2	7.4	7.4	8270	0.33	70 - 130	<u>+</u> 30
	Vinyl chloride	0.2	6	4	8260	0.01	70 – 130	± 30

¹ Analytical methods are from SW-846, "Test Methods for Evaluating Solid Waste, Physical/Chemical Methods," (EPA 1997, as amended).

² The TSCA action limit is 50 ppm total Aroclors for soils. However, ERDF may accept higher PCB concentrations for solid materials.

³ The total halogenated organic carbon action level is from the ERDF WAC. This is calculated from the halogenated organics measured by SW-846 Methods 8260 and 8270.

⁴ The individual radionuclide limits are from Table 3 of the ERDF WAC. They have been converted from Ci/m³ to pCi/g by assuming that the bulk density of the waste is 1.6 g/cm³. Where there are two or more radionuclides present in the waste, the "sum of fractions" method (10 CFR 61.55) shall be used to determine acceptability. This will ensure that the integrated inventory is below TRU and class C limits.

⁵ Total metals and organics will be determined. This is allowed by the regulations (see Method 1311) and will facilitate a rapid turnaround time for analysis. The resulting totals (in mg/Kg) can be compared to the "Action Level for total analysis" in this table or, if applicable, the total can be divided by 20 and compared to the Action Level in the TCLP extract (in mg/L). The factor of 20 is derived from the leachate to sample ratio in the TCLP Method 1311.

⁶ These values represent either the TC action level or the LDR action level, whichever is the lowest concentration.

⁷ ERDF WAC (BHI 1998) for total cresol.

2.3.8 Radiological Field Survey QC

Daily calibration checks will be performed and documented on each survey instrument. Instrument response checks will be made with a known standard in accordance with FH procedures.

2.3.9 Field and Laboratory QC Requirements for Soil Samples

Definitions of these are presented in HASQARD (DOE-RL 1998) and Chapter 1 of SW-846 (EPA 1997). Control samples for field work may include:

- Field equipment blanks will be collected to assess the cleanliness of the sampling equipment and only apply if equipment is reused between events. Therefore, if applicable, equipment blanks will be collected in the field using clean silica sand passed through decontaminated sampling equipment prior to reuse of the equipment. The blank will be analyzed for the same radionuclides and chemical analytes as actual samples collected during the use of the equipment. At least one equipment blank will be collected during sampling for each type of equipment.
- Field duplicates are two samples of the same matrix being sampled. Field duplicates are analyzed independently and provide information concerning the homogeneity of the matrix. At least one field duplicate for each analysis will be submitted.

Control measures taken to monitor laboratory performance are:

- One laboratory method blank for every 20 samples (5% of samples), analytical batch or sample delivery group (whichever is most frequent) will be carried through the complete sample preparation and analytical procedure. The method blank will be used to document contamination resulting from the analytical process.
- One laboratory control sample (LCS) or blank spike will be performed for every batch of samples for each analytical method criteria to monitor the effectiveness of the sample preparation process. The LCS will be carried through the preparation and analysis procedure. Since soil is the matrix of interest, clean sand or water may be used for the LCS. The results from the analyses are used to assess laboratory performance.
- A matrix spike sample will be prepared and analyzed for every 20 samples (as applicable to method) of the same matrix or sample preparation batch, whichever is most frequent. The matrix spike results are used to document the bias of an analytical process in a given matrix.
- Laboratory duplicates or matrix spike duplicates will be used to assess precision and will be analyzed at the same frequency as the matrix spikes. These will be carried through preparation and analysis.

2.3.10 Instrument/Equipment Testing, Inspection, and Maintenance

Measurement and testing equipment used in the field or in the laboratory that directly affects the quality of analytical data will be subject to preventative maintenance measures that ensure minimization of measurement system downtime and avoids inconsistencies in instrument performance.

Laboratories and onsite measurement organizations must maintain their equipment. Instrument preventative maintenance consists of routine inspections, instrument maintenance, and corrective actions. Preventative maintenance is performed in accordance with a schedule based on manufacturer's recommendations, instrument performance history, and usage. Each instrument has a logbook to record maintenance events with date and name of person performing the maintenance. The logbook includes routine inspections, significant corrective actions, instrument maintenance and repairs.

Spare parts inventories help ensure minimal loss of analytical capability. Spare parts include day-to-day consumables and manufactures recommended spare parts.

2.3.11 Instrument Calibration and Frequency

Laboratory measurement systems are subject to calibration and/or calibration verification before use for sample analyses. For all organic laboratory analyses, actual standards of each compound or Aroclors will be used to calibrate the equipment. The number of concentrations of the standards used in the calibration curve will be that required by SW 846 method cited in Table 2-3. Tentatively Identified Compounds analysis cannot be substituted for the use of the actual calibration standards required by the organic method specified in Table 2-3. Calibrations are conducted in accordance with the specific analytical methods performed and in the applicable laboratory QA Plan.

Instruments that fail acceptance criteria shall be investigated and re-calibrated. Instruments are not allowed to be used for sample analysis until they meet acceptance criteria. The responsible chemist or manager is required to take corrective action when measurement systems fail calibration QC criteria.

2.3.12 Inspection/Acceptance Requirements for Supplies and Consumables

FH's procurement activities will be limited to providing BHI Procurement with procurement requisitions. All subject activities will meet the requirements of BHI Procurement procedures found in BHI-PR-01, "ERC Procurement Procedures."

The lot number from the manufacturer-certified, pre-cleaned sample containers is recorded in the sampler's logbook.

2.4 CONTINGENCY ANALYSES

The purpose of contingency sampling and analysis is to verify radionuclide ratios. The purpose of verifying the radionuclide ratios may be to demonstrate that a waste is or is not anomalous. Contingency sampling and analysis will be performed and managed by FH. Determination of anomalous waste is discussed in Section 2.2.3.

2.4.1 When Contingency Analyses May Be Required

Contingency analyses may be required if the measured gamma ratios fall outside the target range of Table 2-2 and the waste is determined to be anomalous as discussed in Section 2.2.3. Contingency analyses also could occur if the waste is designated as potential TRU waste utilizing the weight-to-curie conversion factors previously discussed. Before conducting contingency sampling, K Basin project staff will determine if there are cost-effective alternatives. If contingency sampling and analyses are chosen, a path forward will be developed by EPA, DOE, FH, and ERDF representatives. Sections 2.4.2 through 2.4.9 discuss the anticipated approach to contingency sampling and analyses. The details of the approach may vary depending on the selected vendor and specific waste to be sampled. If conducting a contingency sampling effort, involvement of representatives from ERDF would ensure that the proposed process would provide acceptable data for waste designation.

2.4.2 Contingency Sample Locations, Handling, and Custody Requirements

Waste that has been determined to require sampling will be staged in a controlled area while the path forward is developed. If required, contingency sampling will occur on a representative sample of the waste in the package that is being sampled. The purpose of the contingency sampling is to determine the appropriate representative radionuclide ratios to Cs-137 through radiochemical analysis. It is recommended that beta/gamma and/or alpha survey instruments be used to select a piece of the waste that exhibits a relatively high count rate. This will ensure that adequate contamination is available so the analyses will not be reported as "less-than values."

K Basin operators will be responsible for contingency sample collection, packaging and shipment of samples to the selected laboratory. Any sampling will be conducted in accordance with the guidance presented in HASQARD, Volume II, Section 4.0 and will address the following activities:

- Sample Identification
- Chain of Custody
- Sample Packaging
- Sample Shipment
- Field Logbooks

2.4.3 Contingency Sample Preservation, Containers, Size, and Holding Times

For contingency sampling, preservation is not applicable to the debris or soil samples. Certified clean plastic or glass containers are not necessary for sample collection. Any clean container may be used. It is recommended that at least 500 g of sample be collected in two or more bottles. This will provide a backup sample if needed. The laboratory requires that the waste be cut into pieces of 1 to 2 in² each or less. It is recommended that final sample weight is discussed with the laboratory before obtaining the samples. Holding times for radionuclide analyses are 180 days.

2.4.4 Contingency Sample Shipping

All sample containers will undergo field radiological screening to determine proper shipping and handling requirements. On-site transfers over nonpublic thoroughfares shall be performed in accordance with written procedures. The procedure includes requirements for proper monitoring and control of the radioactive samples and should be reviewed and approved by the Radiological Control Organization. Shipments of waste samples are performed per HNF-PRO-156, "Non-radioactive Hazardous Material/Hazardous Waste Shipments (HM/HW)," if the samples are known or suspected to be mixed waste, and HNF-PRO-157 if the samples are radioactive waste.

2.4.5 Analytical Methods Requirements for Contingency Samples

Parameters and methods for contingency samples are listed in Table 2-3. Laboratory-specific SOPs for analytical methods are in place. Laboratory SOPs and QA plans to be used include analytical procedures and QA plans from 222-S Laboratory. Other laboratories may be used. Changes or additional methods identified during future engineering or planning for contingency sampling should be presented in page changes, addenda, or revisions, as appropriate. Detection limits achievable by the laboratory will be dependent on sample quantity available and also may be affected by the matrix and radionuclide activity levels of the sample.

2.4.6 QC Requirements for Contingency Samples

This characterization effort relies on direct measurements to locate areas of higher beta/gamma contamination for sub-sampling. QA is necessarily built into each phase of the characterization both as QC samples, which monitor sampling and laboratory performance, and field instrument operational checks that monitor field instrumentation performance.

QC measures taken to support field operations performance are described in Section 2.2.6.

For contingency samples collected for laboratory analyses, the following QC samples will be collected during sampling and sent to the laboratory.

Equipment blanks will be collected on contingency samples to assess the potential for gross
cross contamination of the sampling equipment, the effectiveness of the sample
decontamination process, and potential sampling environment contaminant contribution.
These will be used if equipment is not pre-cleaned or if equipment is reused between
sampling events.

Table 2-4. Contingency Sample Measurement Methods, Detection Limits, and Sample Volumes for Selected Radionuclide COCs (2 pages)

Contaminant of Concern (COC)	Analytical Callout	Analytical Technique	Method Reference ¹	Detection Limits ²	Mass Requirements
		1 echnique	Reference	Solid (pCi/g)	Solid (g)
		Radior	iuclides		
H-3 (tritium)	LSC	Liquid Scintillation Beta Analysis	LA-218- 114	400	TBD
Pu-238, Pu-239/240	Pu Isotopic	Alpha Energy Analysis	LA-953- 104	1	TBD
Cm-244	Cm Isotopic	Alpha Energy Analysis	LA-953- 104	1	TBD
Am-241	Am Isotopic	Alpha Energy Analysis	LA-953- 104	1	TBD
Co-60	GEA	Gamma Energy Analysis	LA-548- 121	0.05	TBD
Sb-125	GEA	Gamma Energy Analysis	LA-548- 121	0.5	TBD
Cs-137	GEA	Gamma Energy Analysis	LA-548- 121	1	TBD
Eu-152	GEA	Gamma Energy Analysis	LA-548- 121	0.1	TBD
Eu-154	GEA	Gamma Energy Analysis	LA-548- 121	0.1	TBD
Eu-155	GEA	Gamma Energy Analysis	LA-548- 121	0.1	TBD
Sr-90	Total Radioactive Sr	Beta Counting	LA-220- 103	1	TBD
Ni-63	Sep/LSC	Chemical Separation and Liquid Scintillation Counting	TBD	30	TBD
U-234, U-235, U-238	ICP/MS	ICP/MS	LA-506- 101	1 μg/g	TBD

Table 2-4. Contingency Sample Measurement Methods, Detection Limits, and Sample Volumes for Selected Radionuclide COCs (2 pages)

Contaminant of Concern (COC)	Analytical Callout	Analytical Technique	Method Reference!	Detection Limits? Solid (pCl/g)	Mass Requirements 3 Solid (g)
Pu-238, Pu-239/240	ICP/MS	ICP/MS	LA-506- 101	l μg/g	TBD

¹ An equivalent method may be used dependent on the laboratory performing the analysis.

Control measures taken to monitor laboratory performance are:

- One laboratory method blank for every 20 samples (5% of samples), analytical batch or sample delivery group (whichever is most frequent) will be carried through the complete sample preparation and analytical procedure. The method blank will be used to document contamination resulting from the analytical process.
- One laboratory control sample or blank spike will be performed for every batch of samples
 for each analytical method criteria to monitor the effectiveness of the sample preparation
 process. The results from the analyses are used to assess laboratory performance.
- A matrix spike sample will be prepared and analyzed for every 20 samples (as applicable to method) of the same matrix or sample preparation batch, whichever is most frequent. The matrix spike results are used to document the bias of an analytical process in a given matrix. It is assumed the matrix spike will be added after digestion.
- Laboratory duplicates or matrix spike duplicates will be used to assess precision and will be analyzed at the same frequency as the matrix spikes. Replicate analysis of the etching solution (digestate) of pipe coupons will be used to monitor precision where appropriate.

2.4.7 Instrument/Equipment Testing, Inspection, and Maintenance

See Section 2.3.7 for applicable criteria.

2.4.8 Instrument Calibration and Frequency

See Section 2.3.8 for applicable criteria.

2.4.9 Inspection/Acceptance Requirements for Supplies and Consumables

See Section 2.3.9 for applicable criteria.

² Target detection limits are listed for soil or concrete. The detection limit for actual samples or other materials may be higher.

³ Sample matrix may include 1 to 2 in. sections of metal coupons. The estimated mass for these sections is approximately 80g. Other samples (e.g., concrete, soil) may require different sample sizes and the size of sample collected should be coordinated with the laboratory that will provide the analyses.

2.5 ASSESSMENT/OVERSIGHT FOR SURVEY SAMPLING AND ANALYSIS

QA oversight requirements are described below.

2.5.1 Assessments and Response Actions

Project QA related to FH radiological surveys may conduct random surveillances and assessments in accordance with QA-11-006-02, "Quality Assurance Surveillances," to verify compliance with requirements outlined in this SAP, associated project work packages, procedures, and/or regulatory requirements.

Deficiencies identified during the assessment will be reported in accordance with QA-11-006-02. When necessary, corrective actions will be taken by the Operations Support Manager.

BHI Compliance and Quality Programs group may conduct random surveillance and assessments in accordance with BHI-MA-02, "ERC Project Procedures," and Procedure 2.9, "Surveillances," to verify compliance with the requirements outlined in this SAP, project work packages, the BHI Quality Management Plan, BHI procedures, and regulatory requirements.

Deficiencies identified by one of the assessments shall be reported in accordance with BHI-MS-02, Procedure 5.3, "Self-Assessments." When appropriate corrective actions will be taken by the project engineer in accordance with HASQARD, Volume 1, Section 4.0 (DOE-RL 1996a) to minimize recurrence.

2.5.2 Reports to Management

Management Assessments for FH are performed in accordance with MS-1-036-02, "Management Assessments." Management Assessment results are reviewed and analyzed by management to identify and implement appropriate actions. Management Assessment results are distributed to affected managers and deficiencies and are managed as required by HNF-PRO-052, "Corrective Action Management."

BHI management assessment shall be made aware of all deficiencies identified by the self-assessments. All deficiencies shall be reported in accordance with BHI-MA-02, Procedure 5.3.

2.6 DATA REVIEW, VALIDATION AND USABILITY

Requirements for review and evaluation of data usability are described in the following sections.

2.6.1 Data Review and Verification Requirements

Data verification will be performed on analytical data to assure that sampling and chain-of-custody documentation is complete, sample numbers can be related to the specific sampling location, samples were analyzed within the required holding times, and analyses meet the data quality requirements specified in the SAP.

Analytical personnel and the project team will review the data. Laboratory personnel will perform a peer review of all analytical data. Peer review will be conducted by a person trained to the particular analytical method being reviewed. HASQARD, Volume 4 (DOE-RL 1998) describes the data review that will be performed by the laboratory. The laboratory will use its own data review procedures that meet the HASQARD criteria to review data before it is sent to the project.

Project personnel or designees will review the data and the summary QC with respect to the criteria in this SAP.

Survey measurement systems will be verified by a review of 5% of the documentation to ensure that calibration checks are performed as required by the methods, dates of survey, and analysis locations are properly documented. The review should be performed by program personnel.

2.6.2 Data Validation

Analytical and survey data will not undergo data validation.

2.6.3 Definition of QC Calculations

Following review, the laboratory data will be assessed by the project team against the criteria in Table 2-2. Assessment will include review of quantitative DQO (i.e., accuracy, precision, completeness, and detection limits). These quantitative DQO are defined below.

Precision

If calculated from duplicate measurements:

$$RPD = \frac{(C_1 - C_2)100}{(C_1 + C_2)/2} \tag{1}$$

where:

RPD = relative percent difference

 C_1 = larger of the two observed values

 C_2 = smaller of the two observed values.

If calculated from three or more replicates, use RSD rather than RPD:

$$RSD = \left(s / \overline{y} \right) 100 \tag{2}$$

where:

RSD = relative standard deviation (in units of %)

s = standard deviation

y = mean of replicate analyses.

Standard deviation, s, is defined as follows:

$$s = \sqrt{\sum_{i=1}^{n} \frac{(y_i - \overline{y})^2}{n-1}}$$
(3)

where:

standard deviation

 $\frac{y_I}{y} = \frac{y_I}{y} = \frac{y_I}{y} = \frac{y_I}{y}$ mean of replicate measurements

number of replicates.

Accuracy

For measurements where matrix spikes are used:

$$\%R = 100 \left[\frac{S - U}{C_{sa}} \right] \tag{4}$$

where:

= percent recovery

= measured concentration in spiked aliquot = measured concentration in unspiked aliquot

= actual concentration of spike added.

For situations where a standard reference material (SRM) is used instead of or in addition to matrix spikes:

$$\%R = 100 \left[\frac{C_m}{C_{srm}} \right] \tag{5}$$

where:

= percent recovery

= measured concentration of SRM = actual concentration of SRM.

Completeness

Defined as follows for all measurements:

$$\%C = 100 \left[\frac{V}{T} \right] \tag{6}$$

where:

%C = percent completeness

V = number of measurements judged valid

T = total number of measurements.

Detection Limit

Defined as follows for metals measurements:

$$MDL = t_{(n-1,1-\alpha=0.99)}S \tag{7}$$

where:

MDL = method detection limit

S = standard deviation of the replicate analyses

 $t_{(n-1, 1-\alpha=0.99)}$ = students' t-value appropriate to a 99% confidence level

and a standard deviation estimate with n-1 degree of freedom

2.7 DATA QUALITY ASSESSMENT

Data quality assessment is performed by the project (or project designee) after review of the radiation survey and laboratory data as described in Section 2.6. The review by the project (or project designee) includes evaluation of the method accuracy, precision, detection limits, and completeness as required in Sections 2.2.5, 2.3.8, 2.3.9, 2.4.5, 2.4.6, 3.3 and 3.9.

The data reported must be reviewed with respect to the DQO. This includes the conceptual model and any assumptions that are included in the data collection design. The estimated concentrations of COCs will be compared by the project to the applicable ERDF WAC requirements (BHI 1998) for designation.

2.8 ANALYTICAL DATA REPORTS

The data report required by this SAP is a summary report with QC summary. This report includes a case narrative and analytical QC, such as percent recovery on laboratory control samples, matrix spikes, RPDs on duplicate or matrix spike/matrix spike duplicates, and method blank results and sample result and reporting limits.

3.0 FIELD SURVEY AND SAMPLING OBJECTIVES

3.1 OVERVIEW AND OBJECTIVES

The objective is to characterize the concrete debris, soil, carbon steel debris, and Transite siding debris waste streams to obtain data for designation and disposal. ERDF is the preferred disposal location.

FH will perform the radiological surveys of the packages containing concrete debris, and carbon steel debris and of the Transite walls before demolition. FH will perform the surface soil surveys after the concrete pad is removed. BHI will perform the soil sampling and will submit the soil samples for laboratory analyses. BHI will request routine turn around times for the analyses.

The following provides an overview of the characterization strategy:

- Concrete pad: The pad will be packaged and the weight-to-curie ratio described in Section 2.2.1, and Tables 2-1 and 2-2 will be used, along with the radionuclide ratios based on above water contamination approaches discussed in HNF 2001 in Table 2-2, to provide the radionuclide concentration. Note that the paint used to fix radionuclide contamination on the surface of the concrete will be evaluated for characteristics via the MSDSs in Appendix B. No characteristic metals or organics are listed in the MSDSs.
- Carbon steel monorail and concrete supports: The same approach discussed for the concrete pad will be used for determining the radionuclide content in the steel and the supports. The previous SAP (HNF 2001) and the profile generated based on the previous SAP (ERDF Profile #KBASIN001, Rev 01, 7/19/2001) listed metals in the paint that are on the TC list. The concentration of the metals will be calculated based on the total weight of the debris. The PCB concentration in the paint will be based on previous paint data (FDI 1997).
- Transite wall: The Transite siding is known to contain asbestos and will be managed as such. If the paint appears to be the same older paint used in the monorail, it will be managed in the same manner; otherwise, if the paint is newer (post 1990) the Appendix B MSDSs will be used to profile the waste stream.
- Soil under and around/in front of the pad: Because the basin water may have touched the soil, the radionuclide ratios in Table 2-2 will be verified by analyses, and if verified will be used. Samples for radionuclide analysis will be collected from three hot spots determined by dose rate surveys. There are three options discussed for the timing of soil sampling: (1) sample before digging, (2) sample during the soil removal, and (3) sample from the bags after removal. The timing of sampling is important because the project must remove soil before fixed laboratory data will be completed and thus before a profile and ERDF roll-off (or other approved facility transportation boxes) will be obtained. The project will store soil in large plastic disposal bags until the soil is disposed at ERDF or other approved facility.

For options 1 and 2 for rad, one sample from each of three locations with higher survey data will be composited from the surface to a maximum of 4.5 ft. The maximum sampling depth will be dependent on the planned depth of excavation, which will be provided by FH Project Management. For option 3, the high rad survey areas will be bagged separately and three samples from these 'higher rad survey soil' will be collected.

For all options for rad analysis, the following will be done. Initially GEA of the samples for gamma emitters will be performed by the fixed laboratory. The ratios of individual gamma emitters to Cs-137 will be compared to the values in Table 2-2. If the ratios are the less than or equal to those in Table 2-2, no additional analyses will be performed and the ratios based on Table 2-2 will be used to estimate radionuclides other than Cs-137. If the gamma analysis indicates that the ratios are not consistent with the values in Table 2-2, then radionuclides in Table 2-3 may be analyzed and the actual concentrations used for all measured isotopes.

For options 1 and 2, for the non-radiological analytes, a triangular grid covering the entire area will be used. One sample from each of three random locations will be composited from the surface to a maximum depth of 4.5 ft will be collected. The maximum sampling depth will be dependent on the planned depth of excavation, which will be provided by FH Project Management.

For all three options, all non-rad analyses will be the same and will be those analytes, methods, and PQLs listed in Table 2-3.

Option 3 will be used if the samplers cannot be present during digging and the soil must be bagged before sampling. Triangular grids will be placed on the soil surface after removal of the pad. The grids will cover the soil under the pad and area outside the pad that will be removed. For radionuclides, the surface soil will be surveyed using same grids previously discussed. The three highest dose rate areas will be located and marked with flags. As excavation proceeds, for radionuclides, the soil from the three highest surface survey areas will be placed in one or more bags and will be marked for radionuclide sampling. Three samples will be obtained from the bags containing the soil with high survey data.

For non-rad, the same bags will be randomly numbered, and one sample from each of three randomly chosen bags will be obtained and analyzed as previously discussed.

Workers removing the soil will keep a map and log identifying the location and depth from which each bag of soil is collected. Each bag will be randomly numbered, three random numbers will be selected for sampling the non-radionuclide analytes. A clean, small shovel or trowel or small hand coring device will be used to get samples from various soil portions within each bag. A sample will be collected from each of the three randomly numbered bags.

3.2 RADIOLOGICAL SURVEY OBJECTIVES

This section builds on the DQO Process developed previously (HNF 2001) and summarized in Section 1.0. The sections below summarize the radiological survey and sample design discussed in previous sections. The project objective is to remove all of the debris (e.g., pipe hangers, fuel

storage canisters, miscellaneous tools, hoses) from the K East and K West Basins. The material removed will be washed to remove adhering sludge and disposed as waste debris. Waste from above the basin water line (e.g., protective clothing, cloth, light metal, concrete, ceramic, brick) will also be generated.

Radiological measurements will be utilized to ensure that exposure to radiation and/or radioactive contamination is ALARA. Surveys will ensure that the radiation dose rates and contamination levels are acceptable to the ERDF criteria as specified in the ERDF WAC.

Radiological measurements can be utilized for those wastes that are determined to be anomalous. Those wastes where it is not appropriate to utilize the weight-to-curie relationships due to the density of the waste being outside the range of 0.2 to 8 g/cm³.

Radiological measurements will be provided and can be utilized with sample data and modeling to determine the expected radionuclide concentrations. Determination of the Cs-137 concentration will allow the use of scaling factors identified in Table 2-2 to determine the hard-to-detect radionuclides.

If modeling or direct assay is utilized to make the radionuclide determinations, the modeling methodology or assay results will be provided with the waste package when offered to ERDF personnel for disposal.

3.3 SURVEY LOCATIONS AND FREQUENCY FOR SOIL

The soil under the concrete pad will be surveyed using a dose meter. The surveys will be conducted in accordance with procedures listed in Section 2.2.

3.4 RADIOLOGICAL SURVEY QC.

Radiological survey QC will consist of initial calibrations and operational checks in accordance with the applicable procedures discussed in Section 2.2.5.

3.5 RADIOLOGICAL SAMPLING OBJECTIVES AND REQUIREMENTS

The objective of the soil sampling, as discussed in Section 2.3, is to provide data for use in characterizing the soil for designation and disposal.

The objective of radiological contingency sampling for this project is to provide data to confirm (or establish appropriate) radionuclide ratios for anomalous waste as discussed in Section 2.2.3, Anomalous Waste. Contingency sampling also may be employed to more accurately characterize suspect TRU waste as discussed in Section 2.4.1.

After performing the surveys, the survey reports/maps and locations will be provided to BHI from FH. The radiological control technicians will flag the highest survey locations so that BHI will know where to collect the radiological samples.

Table 3-1. Radiological Survey Instrumentation QC Requirements (2 pages)

				Preliminary		Requirements	
Data Type	Survey Method and Purpose	Analyte	Typical Instrument	Action Level	Detection Limit	Accuracy (% of True Value) (a)	Precision (RSD as %) ^(b)
Dose Rate	Dose rate Measurement for R/hr to Curie Cs-137 conversion and for determination of restricted and nonrestricted waste classification.	Gamma-emitting radionuclides (c)	Eberline RO-3B, Ionization Chamber	50 mR/hr @ 30 cm from surface: 75 mR/hr at surface	0.5 mR/hr	Within Limits printed on source check assembly.	20%
Alpha - Activity	Alpha Scintillation for determination restricted and non restricted waste	Alpha-emitting radionuclides	Bicron Surveyor X with a Scintillation Detector	Fixed Activity: 80,000 dpm/100 cm ² Smears: 400 dpm/100 cm ²	Fixed Activity: <80,000 dpm/100 cm ² Smears: <400 dpm/100 cm ²	Within Limits printed on source check assembly.	20%
Beta/gamma activity	Beta/gamma pancake Geiger-Mueller (GM) for determination restricted and non restricted waste	Beta-emitting radionuclides (d)	Bicron Surveyor X, or Eberline E-140 Series with a pancake GM detector.	Fixed Activity 80,000 dpm/100 cm ² Smears: 100,000 dpm/100 cm ²	Fixed Activity <80,000 dpm/100 cm ² Smears: <100,000 dpm/100 cm ²	Within Limits printed on source check assembly.	20%
Gamma activity	NDA Gamma analysis for determination of radionuclide content of waste.	Gamma-emitting radionuclides	Collimated Gamma Detector, multi- channel analyzer.	45 nCi/g Cs-137 ^(e)	< 45 nCi/g Cs-137	80-120	20%

Table 3-1. Radiological Survey Instrumentation QC Requirements (2 pages)

				Preliminary		Requirements	
Data Type	Survey Method and Purpose	Analyte	Typical Instrument	Action Level	Detection Limit	Accuracy (% of True Value) ^(a)	Precision (RSD as %) ^(b)
Neutron activity	NDA Thermal Neutron analysis for determination of TRU radionuclides.	TRU Radionuclides	Collimated neutron detector	100 nC/g TRU	TBD ^(f)	80-120	20%

^a Source check must be within these limits per applicable procedure.

^b Multiple source checks must within 20% of each other.

Although the instrument is capable of measuring the dose from a wide variety of gamma and beta emitting radionuclides, for purposes of this SAP, the measurements will be made with the window closed and all of the dose will be ascribed to Cs-137.

Although the instrument is capable of measuring gamma emitters with a very low efficiency the response of the instrument will be assumed to be entirely from beta emitting radionuclides.

^e If the waste is such that the radionuclide ratios for K East Basin above water waste are applied, the estimated TRU content of the waste is about 0.4 times the measured Cs-137 activity. Thus, if the method can detect 45 nCi/g Cs-137, then the estimated TRU content would be about 20 nCi/g.

f Acceptable detection limit for neutrons will be such that the detection limit of TRU in waste is equivalent to <50 nCi/g TRU based on estimated TRU content of K East sludge or fuel as appropriate to the waste being measured.

3.6 SOIL SAMPLE COLLECTION

After the concrete pad is removed and the radiological surveys are performed, soil sampling will be performed. Samples will either be collected before soil removal (Option 1), during soil removal (Option 2) or from bags after removal of the soil (Option 3). Two designs are presented: one for radionuclides and another for non-radionuclides.

The sample locations of the three highest rad-surveys will be flagged by rad control technician staff and will be used to sample for SFL analyses for radionuclides. The rad surveys will utilize the same grid established for non-radiological sampling. For options 1 and 2, either a hand auger, shovels, backhoe or bulldozer will be used to collect the sample. A backhoe or bulldozer is allowed because the rock content is not known under the concrete pad and the depth is lower than most hand operated equipment can attain. At each of the three locations, a sample from the surface to 4.5 ft maximum depth will be collected. The sampling crew will be told by the FH Project Manager the depth of excavation in each location. This is done because the depth will vary from a few inches to 4.5 ft, depending on the location. For option 3, the highest rad survey soil will be placed in separate bags and marked as such. Three soil samples will be collected from these 'higher' level bags. Each soil sample for radionuclides at each location will be homogenized in a stainless steel container and will be bottled and shipped to the SFL.

For options 1 and 2, for the non-radioactive analytes, a triangular grid covering the removal area will be generated, each grid node will be numbered and one sample will be collected from each of the three nodes, as identified by the three random numbers. At each of the sample locations, a sample which represents the interval from the surface to depth of excavation (i.e., a few inches to 4.5 ft as provided by FH Project Management) will be taken. Each sample will be composited from the surface to the depth of excavation.

For option 3, bags will be randomly numbered and a sample will be collected from each of three randomly numbered bags. For all options, analysis will be performed as presented in Section 3.1.

For all options, samples for volatile organic analyses will be collected before homogenizing. Each soil sample for metals and all non-volatile organic analyses will be homogenized in a stainless steel container and will be bottled and shipped to the offsite laboratory by the ERC sampling personnel.

The samplers log book should document the following:

- the grid design as applicable,
- the number of any bags and which bags contained the 'higher rad survey soil,'
- location of samples collected and corresponding sample numbers,
- radiological survey results related to the sample locations,
- description of the field conditions and observations.

The soil samples will be obtained with stainless-steel hand augers, shovels, backhoes, or buildozers as tools. Plastic cannot be used for sampling due to the organics being analyzed. It is understood that due to the plastic bags used for storage, phthalates may be present from the storage bags.

3.7 SOIL SAMPLE COLLECTION PROCEDURES

The following procedures found in BHI-EE-01 are used during sample collection:

- Procedure 1.5, "Field Logbooks"
- Procedure 2.0, "Sample Event Coordination"
- Procedure 2.1, "Sampling Documentation Processing"
- Procedure 3.0, "Chain of Custody"
- Procedure 3.1, "Sample Packaging and Shipping"
- Procedure 3.2, "Field Decontamination of Sampling Equipment"
- Procedure 4.0, "Soil and Sediment Sampling"
- Procedure 4.2, "Sample Storage and Shipping Facility"
- Procedure 4.5, "Sample Compositing"

3.8 FIELD AND LABORATORY QC FOR SOIL SAMPLING

Field QC sampling requirements are summarized in Table 3-2. Table 3-3 summarizes the laboratory analytical QC sampling requirements. The definitions of the field and laboratory QC sample are found in Section 2.3.9 of this SAP.

Table 3-2. Field QC Samples and Frequencies, Analytical Methods and Analytes

Analytes and Methods	Field Duplicates	Equipment Blanks
See Table 2-3	Minimum of one. (Normal	Minimum of one. (If
	frequency is 1 per 20	disposable or single-use
	samples, however only 3	equipment is used, no
	samples are specified.)	equipment blanks are
		needed.)

Table 3-3. Laboratory QC Sampling Requirements Summary

💘 QC Sample Type 🤍	Purpose	Frequency Reference
Method blank	Assess laboratory contamination	One per 20 samples of same matrix prepared in one batch.
Laboratory Control Sample	Assess laboratory accuracy	One per 20 samples of same matrix prepared in one batch.
Matrix Spike	Assess accuracy of method on soil matrix	One per 20 samples of same matrix prepared in one batch.
Matrix spike duplicate or duplicate	Assess precision	One per 20 samples of same matrix prepared in one batch.

3.9 SAMPLE MANAGEMENT

3.9.1 Sample Custody

3.9.1.1 Field Custody

All samples obtained during the project will be controlled from the point of origin to the analytical laboratory, as required by BHI-EE-01, Procedure 3.0, "Chain of Custody."

3.9.1.2 Laboratory Custody Procedures

Sample custody during laboratory analysis will be addressed in the applicable laboratory SOPs. Laboratory custody procedures will ensure that sample integrity and identification are maintained through out the analytical process.

3.9.2 Sample Preservation, Containers, and Holding Times for Soils

Soils are preserved via cold storage to slow the chemical reaction rates. Sample preservation and container details will be addressed on the sample authorization form in accordance with BHI-EE-01, Procedure 2.0, "Sample Event Coordination."

Table 3-4.	Soil Sample	e Preservation.	. Minimum	Weight, Containers.	, Storage and Holding Times

Method/Parameter Group	Minimum Collection Weight**	Container P=plastic G=glass	Storage	Holding Time from Collection to Preparation and/or analysis*
6010/metals except Hg	200 g	P or G	Room temperature	6 months from collection to analysis
7471 (Hg)		P or G	4 +/- 2°C	28 days from collection to analysis
8260 Volatiles	20 g	G w/Teflon lid	4 +/- 2°C	14 days from collection to analysis
8270 Semivolatiles, 8151 herbicides, 8081 pesticides 8082 PCBs	500 g	G w/Teflon lid	4 +/- 2°C	14 days from collection to analysis; analyze within 40 days from extraction
GEA	1000g	P or G	Room Temperature	6 months from collection to analysis
AEA/LSC	125 g	P or G	Room Temperature	6 months from collection to analysis

^{*}Turnaround time (TAT) is 45 days for non-radiochemical parameters, and for radiochemical parameters this will be agreed upon by FH and BHI staff.

3.9.3 Sample Packaging and Shipping

Sample packaging and shipping will be performed in accordance with BHI-EE-01, Procedure 3.1, "Sample Packaging and Shipping." Samples submitted for SFL analysis will be screened for radioactivity at the Radiological Counting Facility before shipment offsite.

^{**}The volumes may be altered based on BHI detailed procedures.

4.0 HEALTH AND SAFETY

4.1 GENERAL SAFETY AND HEALTH REQUIREMENTS

All field operations required by this SAP will be conducted in accordance with the HASP (HNF 2000b). Field operation performed by BHI will meet both the FH requirements and the BHI requirements.

4.2 FH SNF REQUIREMETNS

The HASP identifies the primary hazards associated with debris management activities. Some of the hazards included direct radiation exposure, potential personnel contamination, potential inhalation of airborne concentrations of radioactive materials, and exposures to hazardous substances. Rather than list the requirements to mitigate and control radiological and hazardous chemical exposures, the HASP references documents which provide the necessary direction to mitigate and control these hazards. To assist in the development of sub-tier or task-/subproject-specific implementation of the HASP, the Project Hanford Management Contract (PHMC) Automated Job Hazards Analysis (AJHA) will be used in accordance with HNF-PRO-079, "Job Hazard Analysis". The AJHA is a computer-based application to help planners identify the potential hazards associated with a job task, and to implement the proper controls based on the hazards identified. Proper use of the AJHA in conjunction with the project HASP (HNF 2000b), plus specifics associated with the task, will constitute acceptable sub-tier or task-/subproject-specific implementation of the HASP. In accordance with 29 CFR 1910.120(6)(1)(v) (OSHA99A), the HASP (HNF 2000b) shall be made available to PHMC employees and any contractor/subcontractor involved with hazardous waste operations.

The PHMC has a well-developed radiation protection program. This program is described in the HSRCM (DOE-RL 1996). The HSRCM fully implements 10 CFR 835, *Occupational Radiation Protection*, as currently amended. The planning of work involving radiation and radioactive materials hazards is further described in HNF-PRO-1623, "Radiological Work Planning Process." Implementation of radiological work and radiation protection activities is detailed in procedures. Procedures address roles and responsibilities, qualifications, training, implementation of the ALARA philosophy, external and internal dosimetry, monitoring and surveillance, work control mechanisms (e.g., radiation work permits, access and entry requirements), self-assessments, and use of specific radiation monitoring devices and meters.

The PHMC Chemical Management Program (CMP), as described in HNF-PRO-2258 ("Chemical Management"), in conjunction with implementation of the PHMC AJHA in accordance with HNF-PRO-079, will be relied upon to protect the worker, general public, and the environment from specific chemical substances and their associated hazards. The CMP provides direction for the acquisition, storage, transportation, use, final disposition, and record keeping for chemicals at the Hanford Site, as well as management review of the CMP program performance.

4.3 BHI SAFETY REQUIREMENTS

All field operations will be performed in accordance with BHI Health and Safety requirements outline in BHI-SH-01, "ERC Safety and Health Program" and BHI-RC-01, "Radiation Protection Program Manual."

The sampling procedures and associated activities will take into consideration exposure reduction and contamination control techniques that will minimize the radiation exposure to the sampling team as required by BHI-RC-01 and BHI-QA-01, ERC Quality Program.

5.0 REFERENCES

- 10 CFR 61.55, "Waste Classification," Code of Federal Regulations, as amended.
- 10 CFR 835, "Department of Energy Occupational Radiation Protection," *Code of Federal Regulations*, as amended.
- 29 CFR 1910.120, 1991, "Hazardous Waste Operations and Emergency Response," *Code of Federal Regulations*, as amended.
- 40 CFR 268.48, "Land Disposal Restrictions," Code of Federal Regulations, as amended.
- ANSI, 1991, Calibration and Use of Germanium Spectrometers for the Measurement of Gamma-Ray Emission Rates of Radionuclides, ANSI N42.14-1991, American National Standards Institute, New York.
- BHI, 1998, Environmental Restoration Disposal Facility Waste Acceptance Criteria, BHI-00139, Rev. 3, Bechtel Hanford, Inc., Richland, Washington.
- Comprehensive Environmental Response, Compensation, and Liability Act of 1980, 42 USC 9601 et seq.
- DOE-RL, 1996, *Hanford Site Radiological Control Manual*, HSRCM-1, as amended, Prepared for the U.S. Department of Energy, Richland Operations Office by the Hanford Contractors, Richland, Washington.
- DOE-RL, 1998, Hanford Analytical Services Quality Assurance Requirements Documents, DOE/RL-96-68, Rev. 2, U.S. Department of Energy, Richland Operations Office, Richland, Washington.
- DOE-RL, 1999, Focused Feasibility Study for the K Basins Interim Remedial Action, DOE/RL-98-66, Rev. 4, U.S. Department of Energy, Richland Operations Office, Richland, Washington.
- EPA, 1997, Test Methods for Evaluating Solid, Waste Physical/Chemical Methods, SW-846, 3rd Edition, as amended by Updates I (July, 1992), IIA (August, 1993), IIB (January, 1995), and III (1997), U.S. Environmental Protection Agency, Washington, D.C.
- EPA 1999, Interim Action Record of Decision for the 100-BC-1, 100-BC-2 100-DR-1, 100-DR-2, 100-FR-1, 100-FR-2, 100-HR-1, 100-HR-2, 100-KR-1, 100-KR-2, 100-IU-2, 100-IU-6, and 200-CW-3 Operable Units, Hanford Site, Benton County, Washington.
- EPA, DOE, and Ecology, 1999, Declaration of the Record of Decision for DOE Hanford 100 Area, 100-KR-2 Operable Unit, U.S. Department of Energy, U.S. Environmental Protection Agency (EPA 541-R99-059), and Washington State Department of Ecology, Richland, Washington.

- EQM, 2001, Data Quality Objectives Process in Support of Characterization of Soil and Debris from Removal of Structures External to the 100 K Storage Basins, Rev. 0, Environmental Quality Management, Richland, Washington.
- ERDF Profile #KBASIN001, Rev 01, 7/19/2001, prepared by R. Lipinski.
- HNF, 1997, 105-K Basin Material Design Basis Feed Description for Spent Nuclear Fuel Project Facilities, HNF-SD-SNF-TI-009, Rev. 1, Fluor Hanford, Inc., Richland, Washington.
- HNF, 2000a, Data Quality Objectives Process for Designation of K-Basin Debris, HNF-6273, Rev. 0, Fluor Hanford, Inc., Richland, Washington.
- HNF, 2000b, K Basins Interim Remedial Action Health and Safety Plan, HNF-4747, Rev. 1, Spent Nuclear Fuel Project, Richland, Washington.
- HNF, 2000c, 105-K Basin Material Design Basis Feed Description for Spent Nuclear Fuel Project Facilities, HNF-SD-SNF-TI-009, Rev. 3, Volume 1, "Fuel," Fluor Hanford Company, Richland, Washington.
- HNF, 2000d, 105-K Basin Material Design Basis Feed Description for Spent Nuclear Fuel Project Facilities, HNF-SD-SNF-TI-009, Rev. 3, Volume 2, "Sludge," Fluor Hanford Company, Richland, Washington.
- HNF, 2000e, Sampling and Analysis Plan for K Basins' Debris, HNF-6495, Rev. 0, June 2000, Fluor Hanford, Richland, Washington.
- HNF, 2001, Sampling and Analysis Plan for K Basin's Debris, HNF-6495, Rev. 1, February 2001, Fluor Hanford, Richland, Washington.
- Jochen, R.M., 2000, "Letter of Instruction: CACN No. 111456, Spent Nuclear Fuel Facility Operations K Basins Routine Water Samples", From R. M. Jochen to L. L. Lockrem, dated February 28, 2000, Spent Nuclear Fuel Project, Richland, Washington.
- Markel, L.P., 2000, 222-S Laboratory QA Plan, HNF-SD-CP-QAPP-016, Rev. 4, Fluor Hanford, Richland, Washington.
- NRC, 1991, Passive Nondestructive Assay of Nuclear Materials, NUREG/CR-5550, LA-UR-90-732, U.S. Nuclear Regulatory Commission, Washington, D.C.
- Resource Conservation and Recovery Act of 1976, 42 U.S.C 6901 et seq.
- SNF, 2001, Documentation of K Basins Waste Determination Based on Cesium 137 Concentrations in Ci/Kg," SNF-7895, Rev. 0, Fluor Hanford, Richland, Washington.
- Toxic Substances Control Act of 1976, 15 U.S.C., et seq.

- WAC-173-303, Dangerous Waste Regulations, Washington Administrative Code, as amended.
- WHC, 1990, Characterization of Radioactive Waste at 100 Area, WHC-SD-NR-RPT-005, Rev. 0, Westinghouse Hanford Company, Richland, Washington.
- WHC, 1996a, Basis for Dose Rate to Curie Assay Method, WHC-SD-WM-RPT-267, Rev. 0, Westinghouse Hanford Company, Richland, Washington.
- WHC, 1996b, *Procedure for Categorizing and Inventorying Waste in Standard Containers*, WHC-SD-WM-PROC-020, Rev. 0, Westinghouse Hanford Company, Richland, Washington.
- WHC, 1996c, Characterization of Empty Fuel Storage Canisters in 105 KE Basin, WHC-SD-SNF-TI-019, Rev. 0, Westinghouse Hanford Company, Richland, Washington.
- WHC, 1996d, "Analytical Report for K Basin Pipe FT6021", 75745-FAST-96-050, Internal Memo from L. L. Lockrem to R. M. Jochen, dated June 5, 1996, Westinghouse Hanford Company, Richland, Washington.
- WHC, 1996e, Procedure for Categorizing and Inventorying Waste in Standard Containers, WHC-SD-WM-PROC-020, Rev. 0, Westinghouse Hanford Company, Richland, Washington.
- WHC, 1997, Characterization Plan for Spent KE Basin Ion Exchange Modules, WH-SD-SNF-TI-039, Rev. 1, Westinghouse Hanford Company, Richland, Washington.

5.1 SPENT NUCLEAR FUELS PROCEDURES

- AD-14-004, "Radiological Area Access Control."
- MS-1-036-02, "Management Assessments," Effective Date: July 8, 1999, Spent Nuclear Fuel Project Administrative Procedure.
- OP-02-025, "Basin Water Quality Control," Spent Nuclear Fuels Operations Project Technical Procedure.
- OP-43-005E, "Collect Routine Water Samples at 105-KE," Rev. 3D, Issue Date: October 1, 1996, Spent Nuclear Fuels Operations Project Technical Procedure.
- OP-43-006W, "Collect Routine Water Samples at 105-KW," Rev. 4C, Issue Date: February 18, 1999, Spent Nuclear Fuels Operations Project Technical Procedure.
- OP-43-028, "Collect Monthly Center of Basin Water Samples," Rev. 0D, Issue Date: December 9, 1996, Spent Nuclear Fuels Operations Project Technical Procedure.

- OP-43-030, "Transport Water Samples to 222-S Lab," Rev. 0B, Issue Date: December 9, 1996, Spent Nuclear Fuels Operations Project Technical Procedure.
- OP-46-006, "Processing Contaminated Waste for ERDF Disposal," Rev. 0/A, Issue Date: December 13, 1999, Spent Nuclear Fuels Operations Project Technical Procedure.
- QA-11-006-02, "Quality Assurance Surveillances," Effective Date: October 4, 1999, Spent Nuclear Fuel Project Administrative Procedure.
- TN 8-001-08, "General Training Administration," Spent Nuclear Fuel Project Administrative Procedure, Effective Date: February 11, 2000.

5.2 PROJECT HANFORD MANGEMENT SYSTEM PROCEDURES

- HNF-PRO-052, "Corrective Action Management," Rev. 4, Effective Date: April 15, 2000.
- HNF-PRO-079, "Job Hazard Analysis," Rev. 5, Effective Date: May 5, 2000.
- HNF-PRO-156, "Non-radioactive Hazardous Materials/Hazardous Waste (HM/HW) Shipments," Rev. 1, Effective Date: April 15, 2000.
- HNF-PRO-157, "Radioactive Material/Waste Shipments," Rev. 1, Effective Date: April 15, 2000.
- HNF-PRO-338, "Asbestos Control—Construction Industry," Rev. 1, Effective Date: May 23, 2000.
- HNF-PRO-408, "Asbestos—Facility Management/General Industry," Rev. 1, Effective Date: May 18, 2000.
- HNF-PRO-436, "Radiation Protection Instrument Program," Rev. 1, Effective Date: April 15, 2000.
- HNF-PRO-632, "GM Portable Survey Instrument," Rev. 1, Effective Date: March 29, 2000.
- HNF-PRO-633, Rev. 1, "Portable Alpha Meter," Effective Date: April 3, 2000.
- HNF-PRO-648, Rev. 2, "Eberline RO-3B (CP)," Effective Date: March 30, 2000.
- HNF-PRO-1623, Rev. 2, "Radiological Work Planning Process," Effective Date: April 15, 2000.
- HNF-PRO-1892, Rev. 2, "Documentation of Radiological Surveys," Effective Date: April 15, 2000.
- HNF-PRO-2258, Rev. 0, "Chemical Management," Effective Date August 31, 1998.

5.3 WASTE MANGEMENT LABORATORY ANALYTICAL PROCEDURES

- LA-220-103, Rev. F-7, "⁹⁰Strontium in Leachates of Soil, Vegetation, Air Filters and Other Solid Samples," Release Date: November 8, 1999.
- LA-220-104, Rev. E-7, "⁹⁰Strontium in Water by Carbonate Precipitation," Release Date: November 8, 1999.
- LA-506-101, Rev. A-3, "Determination of Trace Elements and Radionuclides by Inductively Coupled Plasma-Mass Spectrometry using TJA Poems," Release Date: November 10, 1999.
- LA-548-121, Rev. F-2, "Preparation of Sample Mounts for Gamma Energy Analysis," Release Date: November 15, 1999.
- LA-953-104, Rev. B-3, "Determination of Plutonium and Americium by Extraction with TRU-SPEC Resin," Release Date: December 21, 1999.

5.4 ERC PROCEDURES

- BHI-EE-01, Environmental Investigations Procedures, Bechtel Hanford, Inc., Richland, Washington.
- BHI-MA-02, ERC Project Procedures, Bechtel Hanford, Inc., Richland, Washington.
- BHI-PR-01, ERC Procurement Procedures, Bechtel Hanford, Inc., Richland, Washington.
- BHI-QA-01, ERC Quality Program, Bechtel Hanford, Inc., Richland, Washington.
- BHI-RC-01, Radiation Protection Program Manual, Bechtel Hanford, Inc., Richland, Washington.
- BHI-SH-01, Hanford ERC Environmental, Safety, and Health Program, Bechtel Hanford, Inc., Richland, Washington.

APPENDIX A INFORMATION SUPPORTING DEVELOPMENT OF RADIONUCLIDE RATIOS FOR CHARACTERIZATION OF K EAST DEBRIS AND SOIL

INFORMATION SUPPORTING DEVELOPMENT OF RADIONUCLIDE RATIOS FOR CHARACTERIZATION OF K East DEBRIS AND SOIL

The source-term for all radionuclides that could reasonably be expected in the 100 K East Area are from N Reactor fuel and associated activation products. The selection of constituents of concern (COCs) was discussed in Appendix A of the Sampling and Analysis Plan for K Basin's Debris, HNF-6495, Rev. 1 (HNF 2001) and Appendix B of the Data Quality Objectives Process for Designation of K-Basin Debris, Rev 0 (HNF 2000a). The selection was performed by listing all radionuclides that have been reported as present in the fuel or measured during historical characterization of the K East, K West, N, or 105-C fuel storage basins. Several selection criteria were applied to define the Environmental Restoration Disposal Facility (ERDF) Waste Acceptance Criteria (WAC) (BHI 1998) that all "Radioactive waste constituents shall be adequately characterized to permit proper segregation, treatment, storage, and/or disposal. This characterization shall ensure that the major radionuclide content of the waste is known and recorded during the waste management process..." (ERDF WAC, Section 3.2.1.1). As a result of that effort, twenty radionuclide COCs were selected. The sections below discuss the application of radionuclide ratios to estimate the radionuclide content of K East Basin debris and soil for those radionuclides that are not measured from radionuclides that are measured.

A.1 RADIONUCLIDE RATIOS FOR CHARACTERIZATION OF K BASIN DEBRIS

Subsequent to the DQO report for the K Basin project (HNF 2000a), an additional two-volume document was obtained. These documents were entitled:

- 1. HNF-SD-SNF-TI-009, Rev 3, 105-K Basin Material Design Basins Feed Description for Spent Nuclear Fuel Project Facilities, Volume 1, "Fuel" (HNF 2000b) and
- 2. HNF-SD-SNF-TI-009, Rev 3, 105-K Basin Material Design Basins Feed Description for Spent Nuclear Fuel Project Facilities, Volume 1, "Sludge" (HNF 2000c).

These two documents formed the basis for the selection of radionuclide ratios for the purpose of estimating the radionuclide content of above water waste from the K Basins. A WHC report (WHC 1990) contained extensive analyses of samples from the K East and K West Basin areas above the water line. These data provided valuable estimates of several radionuclides that had not been estimated from other sources (e.g., nickel-59, chromium-51, and manganese-54). In order to put all radionuclides from the various sources on a normalized basis, all final estimates of radionuclide content of the fuel (HNF 2000b, HNF 2000c) or samples from K West and K East Basins, were converted to a percent of the estimated Cs-137 concentration. For instance, if the reference indicated that the fuel would contain 500 Ci of strontium-90 and 1,000 Ci of Cs-137, the percentage entered into Table A-1 would be 50%. Only the information related to this K East removal project is included in the remainder of this appendix.

In addition to the reports mentioned above there were several sampling efforts that had been conducted on various waste streams. The data from these various sampling efforts were tabulated and reviewed and ratios of each radionuclide measured are presented in Table A-1. Based on a review of the data from the various sources and the conceptual model for the waste

stream it was determined that the following logic would be used to select the applicable ratio for each waste stream. The K East basin could have different sets of ratios that could be applied to the waste depending on the origin of the waste. The ratio applicable to waste that originates from above the water line of the basin was considered applicable to this removal action. The above water waste around the K-Basin was contaminated mainly by water removed from the basins and incidental sludge particles that might have adhered to equipment being moved in and out of the basins. Filters from the basin were placed on the Filter Wash Pad that is being removed. Basin water and some sludge particles would likely be the source of contamination on the wash pad and surrounding soil. Data that measured the airborne contamination radionuclide ratios were also used in developing the K East Basin above water radionuclide ratios, thus the ratios are also applicable to incidental contamination that may be found on the Monorail Foundations/Supports and Rails. The ratios used for this waste are primarily an amalgamation of data from WHC (1990) and data from recent air sampling data (Slotemaker 1999).

Additional discussion regarding the selection of applicable radionuclide ratios is provided below.

A.1.1 Above-Water Waste

Significant differences from radionuclide ratios found in fuel and found in K East versus K West data were noted in historical analyses of samples from above water portions of the K East and K West (WHC 1990). Another source of data that was used was the air sampling data from 1998 (Slotemaker 1999). It was reasoned that the data obtained from collecting high volume air samples would provide a reasonable estimate of the radionuclide mix that might be encountered from continued operations around the fuel basins. In selecting the final ratios to use, generally the highest ratio from either the WHC (1990) report or the air data were used if available for a specific radionuclide. Not all of the COC radionuclides were measured on the samples from either source. If there were no measured ratios, then K East fuel data radionuclide ratios (HNF 2001) were selected. Table A-2 provides a summary of the final selected radionuclide ratios.

A.1.2 Estimate of Gamma Survey Ratios to Define Anomalous Waste

The purpose of estimating a ratio of all COC gamma emitters to Cs-137 is to alert project staff to the presence of waste that is outside of the anticipated ratios that were discussed above. It is assumed that if the measured contamination levels of the six major anticipated gamma emitters (including Cs-137) to the measured Cs-137 activity (dpm/package) is within a certain range, then the waste is presumed to contain contamination that can be adequately estimated using the listed ratios in Table A-2. If the gamma data ratio is outside of the estimated range, then the waste is considered anomalous and will be subjected to additional NDA measurements and/or sampling.

Table A-1. Comprehensive List of Radionuclide Contaminants Concern and Ratios to Cs-137 for K Basin Waste.

Radi	onuclide Symbol	2000 Fuel K East Ratio % to Cs-137 ⁽²⁾	WHC Report K East Ratio % to Cs-137 ⁽¹⁾	K East Air Data % of Cs- 137 ⁽²⁾	Chosen Ratio for K East Above Water Debris		
Tritium	H-3	0.26%	0.09%	2017131111111111111111111111111111111111	0.090%		
Cobalt	Co-60	0.023%	0.96%		1.0%		
Nickel	Ni-63	0.036%	0.34%		0.34%		
Strontium	Sr-90	76%	103.01%	51.39%	103%		
Antimony	Sb-125	0.16%			0.16%		
Cesium	Cs/Ba-137m	100%	100%	100%	100%		
Promethium	Pm-147	2.3%			2.3%		
Samarium	Sm-151	1.4%			1.4%		
Europium	Eu-152	0.062%			0.062%		
Europium	Eu-154	0.72%			1.4%(4)		
Europium	Eu-155	0.13%			0.45%	5	
Uranium	U-234	0.0074%	0.027%		0.027%		
Uranium	U-235	0.0003%	0.0046%		0.0046%	2.	
Uranium	U-238	0.0061%	0.02%		0.021%	6	
Plutonium	Pu-238	0.95%	2.07%	2.0%	2.1%		
Plutonium	Pu-239	1.9%	13.20%	11.6%	13%	ŀ	
Plutonium	Pu-240				5.1%(3)		
Plutonium	Pu-241	50%	197.05%		197%		
Americium	Am-241	8.2%	16.71%	6.88%	17%	Y R	
Curium	Cm-244	0.013%			0.013%		

⁽¹⁾ WHC-SD-NR-RPT-005, Rev 0. "Characterization of Radioactive Waste at 100 Area", Nov. 1990. Written by John DeVanney (WHC 1990)

⁽²⁾ Data from Table 3.6, "105-K Basin Material Design Basis Feed Description for Spent Nuclear Fuel Project Facilities, Volume 1, Fuel," HNF-SD-SNF-TI-009, Volume 1, Rev. 3 (HNF 2000b)

⁽³⁾ Because the air data for the measured Pu isotopes showed an increased % relative to Cs-137, a ratio that was measured in K East Basin sludge was applied in this case (Table A-1, HNF 2001).

⁽⁴⁾ A ratio that was measured in K East Basin sludge was applied in this case (Table A-1, HNF 2001).

Table A-2. Summary List of Radionuclide Contaminants of Concern and Ratios to Cs-137 for K East Debris and Soil.

Radionuclide Name	Radionuclide Symbol	Chosen Ratio for K East Debris and Soil
Tritium	H-3	0.090%
Cobalt	Co-60	1.0%
Nickel	Ni-63	0.34%
Strontium	Sr-90	103%
Antimony	Sb-125	0.16%
Cesium	Cs/Ba-137m	100%
Promethium	Pm-147	2.3%
Samarium	Sm-151	1.4%
Europium	Eu-152	0.062%
Europium	Eu-154	1.4%
Europium	Eu-155	0.45%
Uranium	U-234	0.027%
Uranium	U-235	0.0046%
Uranium	U-238	0.021%
Plutonium	Pu-238	2.1%
Plutonium	Pu-239	13%
Plutonium	Pu-240	5.1%
Plutonium	Pu-241	197%
Americium	Am-241	17%
Curium	Cm-244	0.013%

A.3 REFERENCES

- BHI, 1998, Environmental Restoration Disposal Facility Waste Acceptance Criteria, BHI-00139, Rev. 3, Bechtel Hanford, Inc., Richland, Washington.
- HNF, 1998, Hanford Site Solid Waste Acceptance Criteria, HNF-EP-0063-4, Rev. 5, Fluor Hanford, Richland, Washington.
- HNF, 2000a, Data Quality Objectives Process for Designation of K-Basin Debris, HNF-6273, Rev. 0, Fluor Hanford, Inc., Richland, Washington.
- HNF, 2000b, 105-K Basin Material Design Basis Feed Description for Spent Nuclear Fuel Project Facilities, HNF-SD-SNF-TI-009, Rev. 3, Volume 1, "Fuel," Fluor Hanford, Richland, Washington.
- HNF, 2000c, 105-K Basin Material Design Basis Feed Description for Spent Nuclear Fuel Project Facilities, HNF-SD-SNF-TI-009, Rev. 3, Volume 2, "Sludge," Fluor Hanford, Richland, Washington.
- HNF, 2001, Sampling and Analysis Plan for K Basin's Debris, HNF-6495, Rev. 1, Fluor Hanford, Richland, Washington.
- Slotemaker, 1999, "Facility Source Term Report," 99-SNF/CJS-024, Interoffice Correspondence from P.G. Huntley to C. J. Slotemaker, dated March 29, 1999.
- WHC, 1990, Characterization of Radioactive Waste at 100 Area, WHC-SD-NR-RPT-005, Rev. 0, Westinghouse Hanford Company, Richland, Washington. Written by John DeVanney.

APPENDIX B GFMSDSs FOR PAINT ON CONCRETE PAD

MSDS # 057948

MATERIAL SAFETY DATA SHEET

COATINGS AND RESINS GROUP

PPG Industries, Inc.

SECTION 1 - CHEMICAL, PRODUCT, AND COMPANY INFORMATION

PRODUCT CODE/IDENTITY: 97-948

REVISION DATE: 04/13/00 (000) 0814

CUSTOMER PART #/NAME: Not applicable

PRODUCT TRADE NAME: PITT-GUARD ALL WEATHER DTR GRAY COA

CHEMICAL FAMILY: Epoxy

EMERGENCY MEDICAL/SPILL INFO: (304) 843-1300 (U.S.) 91-800-00-214 (MEXICO)

TECHNICAL INFORMATION: 1-800-441-9695

PRODUCT SAFETY/MSDS INFORMATION: 4325 ROSANNA DRIVE, P.O. BOX 9 ALLISON PARK, PA 15101 (412) 492-5555

DATE OF MSDS PREPARATION: 07/31/00

PRIMARY HAZARD WARNING

Flammable. Keep away from heat, sparks, flames, and other sources of ignition. Do not smoke. Extinguish all flames and pilot lights. Turn off stoves, heaters, electrical motors, and other sources of ignition during use and until all vapors/odors are gone. Harmful if swallowed. May cause slight skin irritation. Causes eye irritation. Prolonged or repeated contact may cause an allergic skin reaction. Vapor and/or spray mist may be harmful if inhaled. Vapor irritates eyes, nose, and throat. Sanding and grinding dusts may be harmful if inhaled.

THIS MATERIAL SAFETY DATA SHEET HAS BEEN PREPARED IN ACCORDANCE WITH THE OSHA HAZARD COMMUNICATION STANDARD (29 CFR 1910.1200), THE SUPPLIER NOTIFICATION REQUIREMENTS OF SARA TITLE III, SECTION 313, AND OTHER APPLICABLE RIGHT-TO-KNOW REGULATIONS.

REF		2 - COMPOSITION/INFO	RMATION ON PERCENT	INGREDIENTS CAS NUMBER	CARCINOGEN*
01 <i>L</i> ETHYI	L BENZENE		1 - <5	100-41-4	
02 KTOLUI	ENE		5 - <10	108-88-3	
03 XXLEI	NES		5 - <10	1330-20-7	
04 CARBO	ON BLACK	•	0.1- <1	1333-86-4	I
05 TITA	NIUM DIOXIDE		5 - <10	13463-67-7	
06 QUAR	ΓZ		20- <30	14808-60-7	INO
07 ∠⊊₽OX	Y RESIN		40- <50	25068-38-6	
08 SILIC	CA		1 - <5	7631-86-9	

^{*} Carcinogens: O=OSHA; A=ACGIH; N=NTP; I=IARC

MSDS # 057948

SARA TITLE III & CERCLA CLASSIFICATIONS

				SA	RA	31:	1/3:	12
REF	SARA 102 RQ (LBS)	SARA 302 TPQ (LBS)	SARA 313	AC	CH	FL	PR	RE
01	1000	NOT ESTAB	Y	Y '	Y	Υ	N	N
02	1000	NOT ESTAB	Y	Y	N	Y	N	N
03	100	NOT ESTAB	Y	Y	N	Y	N	N
04	NOT ESTAB	NOT ESTAB	N	И	Y	N	N	N
05	NOT ESTAB	NOT ESTAB	И	N :	N	И	N	N
06	NOT ESTAB	NOT ESTAB	N	N	Y	N	И	N
07	NOT ESTAB	NOT ESTAB	N	Y	N	N	14	N
08	NOT ESTAB	NOT ESTAB	И	N	N	N	И	14

SARA 311/312 CATEGORIES FOR THIS PRODUCT: ACUTE= Y, CHRONIC= Y, FLAMMABILITY= Y, PRESSURE= N, REACTIVITY= N

OCCUPATIONAL EXPOSURE LIMITS HAVE BEEN ESTABLISHED FOR THE FOLLOWING MATERIALS:

		wicer:	F1.	0.5.	03114
REF		TLV-TWA	TLV-STEL	PEL-TWA	PEL-STEL
01		100 ppm	125 ppm	100 ppm	125 ppm
02 03	ş-	50 ppm 100 ppm	NOT ESTAB. 150 ppm	100 ppm 100 ppm	150 ppm 150 ppm
04		3.5 mg/m3	NOT ESTAB.	3.5 mg/m3	NOT ESTAB.
05 06	R-	10 mg/m3 0.1 mg/m3	NOT ESTAB. NOT ESTAB.	10 mg/m3 R- 0.1 mg/m3	NOT ESTAB.
07		NOT ESTAB.	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.
80		10 mg/m3	NOT ESTAB.	6 mg/m3	NOT ESTAB.

[C- Ceiling Limit; S- Potential Skin Absorption; R- Respirable Dust] [NOT ESTAB. = NOT ESTABLISHED = NOT APPLICABLE]

PRODUCT STATUS RELATIVE TO THE U.S. EPA TOXIC SUBSTANCES CONTROL ACT

All chemical substances in this product are listed on the U.S. TSCA Inventory or are otherwise exempt from TSCA Inventory reporting requirements.

SECTION 3 - HAZARDS IDENTIFICATION

EFFECTS OF OVEREXPOSURE FROM:

INGESTION: Harmful if swallowed.

EYE CONTACT: Causes eye irritation.

SKIN CONTACT: May cause slight skin irritation. Prolonged or repeated contact may cause an allergic skin reaction.

INHALATION: Vapor and/or spray mist may be harmful if inhaled. Vapor irritates eyes, nose, and throat. Sanding and grinding dusts may be harmful if inhaled. Repeated exposure to high vapor concentrations may cause irritation of the respiratory system and permanent brain and nervous system damage.

CHRONIC OVEREXPOSURE: Avoid long-term and repeated contact. This product contains crystalline silica which has been classified as a human carcinogen by IARC. Long-term exposures may also lead to a disabling lung condition known as silicosis. The risk depends on the duration and level of exposure to dust from

MSDS # 057948

sanding surfaces or mist from spray applications. Use of appropriate personal protective equipment and/or engineering controls should be employed whenever these types of operations are being performed. This product contains titanium dioxide. Animals inhaling massive quantities of titanium dioxide dust in a long-term study developed lung tumors. Studies with humans involved in manufacture of this pigment indicate no increased risk of cancer from exposure. Potential for inhalation of titanium dioxide dusts from coatings is very limited. Since overexposures are not expected, there is no significant hazard for man. This product contains toluene. Toluene inhalation in animals (greater than 1500 ppm) and intentional inhalation of toluene-containing products by humans (e.g. glue) has caused adverse fetal development effects. This product contains carbon black which has been rated an IARC 2B carcinogen due to animal data. Ethylbenzene has been reported by NTP to cause cancer in laboratory animals following a chronic (2 year) inhalation exposure. Dose levels of 75, 250 and 750 ppm were used, with evidence of carcinogenicity found in the kidneys of rats and the lung and liver of mice at 750 ppm. The No Observed Effect Level (NOEL) was 75 ppm. The relevance of these findings to humans is uncertain, but appropriate safeguards should be employed to reduce or eliminate inhalation exposure to ethylbenzene. High exposures to xylenes in some animal studies have been reported to cause health effects on the developing embryo and fetus. These effects were often at levels toxic to the mother. The significance of these findings to humans has not been determined.

SIGNS AND SYMPTOMS OF OVEREXPOSURE: Eye watering, headaches, nausea, dizziness, and loss of coordination are indications that solvent levels are too high. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal. Redness, itching, burning sensation and visual disturbances may indicate excessive eye contact. Dryness, itching, cracking, burning, redness, and swelling are conditions associated with excessive skin contact.

MEDICAL CONDITIONS AGGRAVATED BY EXPOSURE: Not applicable.

WARNING: This product contains a chemical(s) known to the State of California to cause cancer and birth defects or other reproductive harm.

SECTION 4 - FIRST AID MEASURES

INGESTION: If swallowed, do not induce vomiting. Gently wipe out inside mouth to remove any residual material.

EYE CONTACT: In case of eye contact, remove contact lenses and flush eyes immediately with a gentle stream of luke warm water for at least 15 minutes.

SKIN CONTACT: In case of skin contact, flush immediately with plenty of water for at least 15 minutes followed by washing with soap and water.

INHALATION: If affected by inhalation of vapor or spray mist, remove to fresh air. Apply artificial respiration and other support measures as required.

OTHER: If ingestion, any type of overexposure or symptoms of overexposure occur during or following the use of this product, contact a poison control center, emergency room or physician immediately; have Material Safety Data Sheet information available.

SECTION 5 - FIRE FIGHTING MEASURES

FLASHPOINT: 70 Degrees F' ('21 Degrees C) (PENSKY-MARTENS CLOSED CUP)

MSDS # 057948

FLAMMABLE LIMITS: Lower explosion limit (LEL): 1.2

Upper explosion limit (UEL): Not available

EXTINGUISHING MEDIA: Use National Fire Protection Association (NFPA) Class B extinguishers (carbon dioxide, dry chemical, or universal aqueous film forming foam) designed to extinguish NFPA Class IB flammable liquid fires.

UNUSUAL FIRE AND EXPLOSION HAZARDS: Keep this product away from heat, sparks, flame, and other sources of ignition (i.e., pilot lights, electric motors, static electricity). Invisible vapors can travel to a source of ignition and flash back. Do not smoke while using this product. Keep containers tightly closed when not in use. Closed containers may explode when overheated. Do not apply to hot surfaces. Toxic gases may form when this product comes in contact with extreme heat.

SPECIAL FIRE FIGHTING PROCEDURES: Water spray may be ineffective. Water spray may be used to cool closed containers to prevent pressure build-up and possible autoignition or explosion when exposed to extreme heat. If water is used, fog nozzles are preferable. Fire-fighters should wear self-contained breathing apparatus and full protective clothing.

SECTION 6 - ACCIDENTAL RELEASE MEASURES

STEPS TO BE TAKEN IN CASE MATERIAL IS RELEASED OR SPILLED: Provide maximum ventilation. Only personnel equipped with proper respiratory, skin, and eye protection should be permitted in the area. Remove all sources of ignition. Take up spilled material with sand, vermiculite, or other noncombustible absorbent material and place in clean, empty containers for disposal. Only the spilled material and the absorbant should be placed in this container.

WASTE DISPOSAL METHOD: Waste material must be disposed of in accordance with federal, state, provincial, and local environmental control regulations. Empty containers should be recycled or disposed of through an approved waste management facility.

SECTION 7 - HANDLING AND STORAGE

HANDLING AND STORAGE PRECAUTIONS: Do not store above 120 degrees F. (48 degrees C.). Store large quantities in buildings designed and protected for storage of NFPA Class IB flammable liquids.

OTHER PRECAUTIONS: Vapors may collect in low areas. If this material is part of a multiple component system, read the Material Safety Data Sheet(s) for the other component or components before blending as the resulting mixture may have the hazards of all of its parts. Containers should be grounded when pouring. Avoid free fall of liquids in excess of a few inches.

SECTION 8 - EXPOSURE CONTROLS AND PERSONAL PROTECTION

PERSONAL PROTECTIVE EQUIPMENT FOR:

EYE PROTECTION: Wear chemical-type splash goggles when possibility exists for eye contact due to splashing or spraying liquid, airborne particles, or vapors.

MSDS # 057948

gloves should be constructed of: nitrile rubber. No specific permeation/degradation testing have been done on protective clothing for this product. Recommendations for skin protection are based on infrequent contact with this product. For frequent contact or total immersion, contact a manufacturer of protective clothing for appropriate chemical impervious equipment.

RESPIRATORY PROTECTION: Overexposure to vapors may be prevented by ensuring proper ventilation controls, vapor exhaust or fresh air entry. A NIOSH- approved air purifying respirator with the appropriate chemical cartridges or a positive-pressure, air-supplied respirator may also reduce exposure. Read the respirator manufacturer's instructions and literature carefully to determine the type of airborne contaminants against which the respirator is effective, its limitations, and how it is to be properly fitted and used.

OTHER EQUIPMENT: Clean contaminated clothing and shoes.

VENTILATION REQUIREMENTS: Provide general dilution or local exhaust ventilation in volume and pattern to keep the concentration of ingredients listed in Section 2 below the lowest suggested exposure limits, the LEL below the stated limit, and to remove decomposition products during welding or flame cutting.

SECTION 9 - PHYSICAL AND CHEMICAL PROPERTIES

[FORMULA VALUES, NOT SALES SPECIFICATIONS]

BOILING RANGE: 230- 293Degrees F

SOLUBILITY IN WATER: .0 %,

VAPOR PRESSURE: 13.0 mmHg

WEIGHT/GALLON (LBS): 11.64 (U.S.)

VAPOR DENSITY: Heavier than air

pH: Not applicable

% VOLATILE/VOLUME: 24.630

% SOLIDS BY WEIGHT: 84.73

SPECIFIC GRAVITY: 1.397

EVAPORATION RATE (BuOAc=100): 141

ODOR/APPEARANCE: Viscous liquid with an odor characteristic of the solvents listed in Section 2.

SECTION 10 - STABILITY AND REACTIVITY

This product is normally stable and will not undergo hazardous reactions.

INCOMPATIBILITY (MATERIALS AND CONDITIONS TO AVOID): Avoid contact with strong alkalies, strong mineral acids, or strong oxidizing agents.

HAZARDOUS DECOMPOSITION PRODUCTS: May produce the following hazardous

MSDS # 057948

decomposition products when exposed to extreme heat: carbon monoxide; carbon dioxide; lower molecular weight polymer fractions; Extreme heat includes, but is not limited to, flame cutting, brazing, and welding.

Hazardous Materials Identification System (HMIS) and National Fire Protection Association (NFPA) Ratings:

	NFPA Rating		
2*	HEALTH	2	
3	FLAMMABILITY	3	
0	INSTABILITY	0	
	2*	2* HEALTH 3 FLAMMABILITY	

Rating System: 0=Minimal, 1=Slight, 2=Moderate, 3=Serious, 4=Severe, *=Chronic Effects.

Safe handling of this product requires that all of the information on the MSDS be evaluated for specific work environments and conditions of use.

THIS IS THE END OF THE MSDS FOR: 97-948 (00172120.00197-948)

Manufactured and Supplied by:

PPG INDUSTRIES, EAST POINT

1377 OAKLEIGH DRIVE

EAST POINT, GA 30344

MSDS # 057949

MATERIAL SAFETY DATA SHEET

COATINGS AND RESINS GROUP

PPG Industries, Inc.

SECTION 1 - CHEMICAL, PRODUCT, AND COMPANY INFORMATION

PRODUCT CODE/IDENTITY: 95-249

REVISION DATE: 08/21/98 (000) 0814

CUSTOMER PART #/NAME: Not applicable

PRODUCT TRADE NAME: EPOXY MASTIC CATALYST COMP B

CHEMICAL FAMILY: Epoxy

EMERGENCY MEDICAL/SPILL INFO: (304) 843-1300 (U.S.) 91-800-00-214 (MEXICO)

TECHNICAL INFORMATION: 1-800-441-9695

PRODUCT SAFETY/MSDS INFORMATION: 4325 ROSANNA DRIVE, P.O. BOX 9 ALLISON PARK, PA 15101 (412) 492-5555

DATE OF MSDS PREPARATION: 10/14/99

PRIMARY HAZARD WARNING

Flammable. Keep away from heat, sparks, flames, and other sources of ignition. Do not smoke. Extinguish all flames and pilot lights. Turn off stoves, heaters, electrical motors, and other sources of ignition during use and until all vapors/odors are gone. Harmful if swallowed. May cause moderate skin irritation. Causes eye irritation. May be absorbed through the skin. Prolonged or repeated contact may cause an allergic skin reaction. Vapor and/or spray mist may be harmful if inhaled. May cause irritation and/or allergic respiratory reaction in lungs. Vapor irritates eyes, nose, and throat.

THIS MATERIAL SAFETY DATA SHEET HAS BEEN PREPARED IN ACCORDANCE WITH THE OSHA HAZARD COMMUNICATION STANDARD (29 CFR 1910.1200), THE SUPPLIER NOTIFICATION REQUIREMENTS OF SARA TITLE III, SECTION 313, AND OTHER APPLICABLE RIGHT-TO-KNOW REGULATIONS.

	SECTION 2 - COMPOSITION/IN	FORMATION ON	INGREDIENTS	
REF	HAZARDOUS INGREDIENTS	PERCENT	CAS NUMBER	CARCINOGEN*
	HYL BENZENE		100-41-4	
02 PRO	OPYLENE GLYCOL MONOMETHYL ETHER	5 - <10	107-98-2	
03 ALK	(YL GLYCIDYL ETHER	5 ~ <10	120547-52-6	
04 XXI	LENES	1 - <5	1330-20-7	
05 200	DIUM ALUMINUM SILICATE	20- <30	1344-00-9	
06 JAI	LC C	20- <30	14807-96-6	
07 EPO	DXY RESIN	30- <40	25068-38-6	
08 NIT	TROETHANE	1 ~ <5	79-24-3	

^{*} Carcinogens: O=OSHA; A=AdGIH; N=NTP; I=IARC

MSDS # 057949

SARA TITLE III & CERCLA CLASSIFICATIONS

				SARA	311/312
REF	SARA 102 RQ (LBS)	SARA 302 TPQ (LBS)	SARA 313	AC CH	FL PR RE
01	1000	NOT ESTAB	Y	Y Y	Y N N
02	NOT ESTAB	NOT ESTAB	N	Y N	Y N N
03	NOT ESTAB	NOT ESTAB	N	Y Y	N N N
04	100	NOT ESTAB	Y	A M	Y N N
05	NOT ESTAB	NOT ESTAB	И	YИ	N N N
06	NOT ESTAB	NOT ESTAB	И	n n	N N N
07	NOT ESTAB	NOT ESTAB	N	Y N	N N N
08	NOT ESTAB	NOT ESTAB	N	Y N	Y N N

SARA 311/312 CATEGORIES FOR THIS PRODUCT: ACUTE= Y, CHRONIC= Y, FLAMMABILITY= Y, PRESSURE= N, REACTIVITY= N

OCCUPATIONAL EXPOSURE LIMITS HAVE BEEN ESTABLISHED FOR THE FOLLOWING MATERIALS:

		ACGI	n.	0.5.	USHA
REF		TLV-TWA	TLV-STEL	PEL-TWA	PEL-STEL
01		100 ppm	125 nnm	100	125
02		100 ppm	125 ppm 150 ppm	100 ppm 100 ppm	125 ppm 150 ppm
03		NOT ESTAB	NOT ESTAB	NOT ESTAB	NOT ESTAB
04		100 ppm	150 ppm	100 ppm	150 ppm
05	_	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.
06 07	R-	2 mg/m3 NOT ESTAB.	NOT ESTAB.	R- 2 mg/m3	NOT ESTAB.
08		100 ppm	NOT ESTAB.	NOT ESTAB. 100 ppm	NOT ESTAB.
		55	nor borno.	zoo ppm	HOI DOIND.

[C- Ceiling Limit; S- Potential Skin Absorption; R- Respirable Dust] [NOT ESTAB. = NOT ESTABLISHED = NOT APPLICABLE]

PRODUCT STATUS RELATIVE TO THE U.S. EPA TOXIC SUBSTANCES CONTROL ACT

All chemical substances in this product are listed on the U.S. TSCA Inventory or are otherwise exempt from TSCA Inventory reporting requirements.

SECTION 3 - HAZARDS IDENTIFICATION

EFFECTS OF OVEREXPOSURE FROM:

INGESTION: Harmful if swallowed.

EYE CONTACT: Causes eye irritation.

SKIN CONTACT: May cause moderate skin irritation. May be absorbed through the skin. Prolonged or repeated contact may cause an allergic skin reaction.

INHALATION: Vapor and/or spray mist may be harmful-if inhaled. May cause irritation and/or allergic respiratory reaction in lungs. Vapor irritates eyes, nose, and throat. Repeated exposure to high vapor concentrations may cause irritation of the respiratory system and permanent brain and nervous system damage.

CHRONIC OVEREXPOSURE: Avoid long-term and repeated contact. This product contains nitroethane. Studies with laboratory animals have shown that ingestion

MSDS # 057949

or inhalation of high levels of nitroethane causes kidney and liver damage and central nervous system effects. This product contains talc. In a lifetime inhalation study female rats exposed to an elevated respirable concentration (9 times the Permissible Exposure Limit) of cosmetic grade talc developed lung cancer. To date, no U.S. regulatory agency has classified talc as a carcinogen based on this data. Ethylbenzene has been reported by NTP to cause cancer in laboratory animals following a chronic (2 year) inhalation exposure. Dose levels of 75, 250 and 750 ppm were used, with evidence of carcinogenicity found in the kidneys of rats and the lung and liver of mice at 750 ppm. The No Observed Effect Level (NOEL) was 75 ppm. The relevance of these findings to humans is uncertain, but appropriate safeguards should be employed to reduce or eliminate inhalation exposure to ethylbenzene. High exposures to xylenes in some animal studies have been reported to cause health effects on the developing embryo and fetus. These effects were often at levels toxic to the mother. The significance of these findings to humans has not been determined.

SIGNS AND SYMPTOMS OF OVEREXPOSURE: Eye watering, headaches, nausea, dizziness, and loss of coordination are indications that solvent levels are too high. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal. Redness, itching, burning sensation and visual disturbances may indicate excessive eye contact. Dryness, itching, cracking, burning, redness, and swelling are conditions associated with excessive skin contact.

MEDICAL CONDITIONS AGGRAVATED BY EXPOSURE: Not applicable.

SECTION 4 - FIRST AID MEASURES

INGESTION: If swallowed, do not induce vomiting. Gently wipe out inside mouth to remove any residual material.

EYE CONTACT: In case of eye contact, remove contact lenses and flush eyes immediately with a gentle stream of luke warm water for at least 15 minutes.

SKIN CONTACT: In case of skin contact, flush immediately with plenty of water for at least 15 minutes followed by washing with soap and water.

INHALATION: If affected by inhalation of vapor or spray mist, remove to fresh air. Apply artificial respiration and other support measures as required.

OTHER: If ingestion, any type of overexposure or symptoms of overexposure occur during or following the use of this product, contact a poison control center, emergency room or physician immediately; have Material Safety Data Sheet information available.

SECTION 5 - FIRE FIGHTING MEASURES

FLASHPOINT: 80 Degrees F (26 Degrees C) (PENSKY-MARTENS CLOSED CUP)

FLAMMABLE LIMITS: Lower explosion limit (LEL): 2.1

Upper explosion limit (UEL): Not available

EXTINGUISHING MEDIA: Use National Fire Protection Association (NFPA) Class B extinguishers (carbon dioxide, dry chemical, or universal aqueous film forming foam) designed to extinguish NFPA Class IC flammable liquid fires.

UNUSUAL FIRE AND EXPLOSION HAZARDS: Keep this product away from heat, sparks,

MSDS # 057949

flame, and other sources of ignition (i.e., pilot lights, electric motors, static electricity). Invisible vapors can travel to a source of ignition and flash back. Do not smoke while using this product. Keep containers tightly closed when not in use. Closed containers may explode when overheated. Do not apply to hot surfaces. Toxic gases may form when this product comes in contact with extreme heat.

SPECIAL FIRE FIGHTING PROCEDURES: Water spray may be ineffective. Water spray may be used to cool closed containers to prevent pressure build-up and possible autoignition or explosion when exposed to extreme heat. If water is used, fog nozzles are preferable. Fire-fighters should wear self-contained breathing apparatus and full protective clothing.

SECTION 6 - ACCIDENTAL RELEASE MEASURES

STEPS TO BE TAKEN IN CASE MATERIAL IS RELEASED OR SPILLED: Provide maximum ventilation. Only personnel equipped with proper respiratory, skin, and eye protection should be permitted in the area. Remove all sources of ignition. Take up spilled material with sand, vermiculite, or other noncombustible absorbent material and place in clean, empty containers for disposal. Only the spilled material and the absorbant should be placed in this container.

WASTE DISPOSAL METHOD: Waste material must be disposed of in accordance with federal, state, provincial, and local environmental control regulations. Empty containers should be recycled or disposed of through an approved waste management facility.

SECTION 7 - HANDLING AND STORAGE

HANDLING AND STORAGE PRECAUTIONS: Do not store above 120 degrees F. (48 degrees C.). Store large quantities in buildings designed and protected for storage of NFPA Class IC flammable liquids.

OTHER PRECAUTIONS: Vapors may collect in low areas. If this material is part of a multiple component system, read the Material Safety Data Sheet(s) for the other component or components before blending as the resulting mixture may have the hazards of all of its parts. Containers should be grounded when pouring. Avoid free fall of liquids in excess of a few inches.

SECTION 8 - EXPOSURE CONTROLS AND PERSONAL PROTECTION

PERSONAL PROTECTIVE EQUIPMENT FOR:

EYE PROTECTION: Wear chemical-type splash goggles when possibility exists for eye contact due to splashing or spraying liquid, airborne particles, or vapors.

SKIN PROTECTION: Wear protective clothing to prevent skin contact. Apron and gloves should be constructed of: impermeable material. No specific permeation/degradation testing have been done on protective clothing for this product. Recommendations for skin protection are based on infrequent contact with this product. For frequent contact or total immersion, contact a manufacturer of protective clothing for appropriate chemical impervious equipment.

RESPIRATORY PROTECTION: Overexposure to vapors may be prevented by ensuring proper ventilation controls, vapor exhaust or fresh air entry. A NIOSH- approved air purifying respirator. with the appropriate chemical cartridges or a

MSDS # 057949

positive-pressure, air-supplied respirator may also reduce exposure. Read the respirator manufacturer's instructions and literature carefully to determine the type of airborne contaminants against which the respirator is effective, its limitations, and how it is to be properly fitted and used.

OTHER EQUIPMENT: Clean contaminated clothing and shoes.

VENTILATION REQUIREMENTS: Provide general dilution or local exhaust ventilation in volume and pattern to keep the concentration of ingredients listed in Section 2 below the lowest suggested exposure limits, the LEL below the stated limit, and to remove decomposition products during welding or flame cutting.

SECTION 9 - PHYSICAL AND CHEMICAL PROPERTIES

[FORMULA VALUES, NOT SALES SPECIFICATIONS]

BOILING RANGE: 248- 396Degrees F

SOLUBILITY IN WATER: 8.8 %

VAPOR PRESSURE: 9.2 mmHg

WEIGHT/GALLON (LBS): 12.64 (U.S.)

VAPOR DENSITY: Heavier than air

pH: Not applicable

% VOLATILE/VOLUME: 21.280

% SOLIDS BY WEIGHT: 86.95

SPECIFIC GRAVITY: 1.517

EVAPORATION RATE (BuOAc=100): 56

ODOR/APPEARANCE: Viscous liquid with an odor characteristic of the solvents listed in Section 2.

SECTION 10 - STABILITY AND REACTIVITY

This product is normally stable and will not undergo hazardous reactions.

INCOMPATIBILITY (MATERIALS AND CONDITIONS TO AVOID): Avoid contact with strong alkalies, strong mineral acids, or strong oxidizing agents.

HAZARDOUS DECOMPOSITION PRODUCTS: May produce the following hazardous decomposition products when exposed to extreme heat: carbon monoxide; carbon dioxide; lower molecular weight polymer fractions; Extreme heat includes, but is not limited to, flame cutting, brazing, and welding.

Hazardous Materials Identification System (HMIS) and National Fire Protection Association (NFPA) Ratings:

nmis kating		NFPA Rating	
		~	
HEALTH	3*	HEALTH	3
FLAMMABILITY	3 , '	FLAMMABILITY	3

MSDS # 057949

REACTIVITY 0 INSTABILITY 0

Rating System: 0=Minimal, 1=Slight, 2=Moderate, 3=Serious, 4=Severe, *=Chronic Effects.

Safe handling of this product requires that all of the information on the MSDS be evaluated for specific work environments and conditions of use.

THIS IS THE END OF THE MSDS FOR: 95-249 (00125847.00195-249)

Manufactured and Supplied by:

PPG INDUSTRIES, EAST POINT

1377 OAKLEIGH DRIVE

EAST POINT, GA 30344

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MSDS # 057950

MATERIAL SAFETY DATA SHEET

COATINGS AND RESINS GROUP

PPG Industries, Inc.

SECTION 1 - CHEMICAL, PRODUCT, AND COMPANY INFORMATION

PRODUCT CODE/IDENTITY: 95-2402

REVISION DATE: 09/29/98 (000) 0814

CUSTOMER PART #/NAME: Not applicable

PRODUCT TRADE NAME: PITT-GUARD RAPID CURE YELLOW BASE

CHEMICAL FAMILY: Polyamide

EMERGENCY MEDICAL/SPILL INFO: (304) 843-1300 (U.S.) 91-800-00-214 (MEXICO)

TECHNICAL INFORMATION: 1-800-441-9695

PRODUCT SAFETY/MSDS INFORMATION: 4325 ROSANNA DRIVE, P.O. BOX 9 ALLISON PARK, PA 15101 (412) 492-5555

DATE OF MSDS PREPARATION: 10/14/99

PRIMARY HAZARD WARNING

Flammable. Keep away from heat, sparks, flames, and other sources of ignition. Do not smoke. Extinguish all flames and pilot lights. Turn off stoves, heaters, electrical motors, and other sources of ignition during use and until all vapors/odors are gone. Harmful if swallowed. May cause skin burns. This product contains a material which causes irreversible eye damage. May be absorbed through the skin. Prolonged or repeated contact may cause an allergic skin reaction. Vapor and/or spray mist harmful if inhaled. Vapor irritates eyes, nose, and throat.

THIS MATERIAL SAFETY DATA SHEET HAS BEEN PREPARED IN ACCORDANCE WITH THE OSHA HAZARD COMMUNICATION STANDARD (29 CFR 1910.1200), THE SUPPLIER NOTIFICATION REQUIREMENTS OF SARA TITLE III, SECTION 313, AND OTHER APPLICABLE RIGHT-TO-KNOW REGULATIONS.

	SECTION 2 - COMPOSITION/INFOR	MATION ON	INGREDIENTS	
REF	HAZARDOUS INGREDIENTS	PERCENT	CAS NUMBER	CARCINOGEN*
01	ETHYL BENZENE	1 - <5	100-41-4	
02 ;	BENZYL ALCOHOL	5 - <10	100-51-6	
03	XYLENES	5 - <10	1330-20-7	
04	SODIUM ALUMINUM SILICATE	10- <20	1344-00-9	
05	TALC	5 - <10	14807-96-6	
06	ISOPROPYL ALCOHOL	10- <20	67-63-0	
07	POLYAMIDE RESIN	40- <50	68410-23-1	
08	SUBSTITUTED AMIDE	1 - <5	82199-12-0	
09	2,4,6 TRIS (DIMETHYLAMINOMETHYL) PHENOL	1 - <5	90-72-2	

MSDS # 057950

* Carcinogens: O=OSHA; A=ACGIH; N=NTP; I=IARC

SARA TITLE III & CERCLA CLASSIFICATIONS

				SARA	311/312
REF	SARA 102 RQ (LBS)	SARA 302 TPQ (LBS)	SARA 313	AC CH	FL PR RE
01	1000	NOT ESTAB	Y	Y Y	Y N N
02	NOT ESTAB	NOT ESTAB	И	Y N	Y N N
03	100	NOT ESTAB	Y	Y N	Y N N
04	NOT ESTAB	NOT ESTAB	N	Y N	N N N
05	NOT ESTAB	NOT ESTAB	N	N N	N N N
06	NOT ESTAB	NOT ESTAB	N	A M	Y N N
07	NOT ESTAB	NOT ESTAB	N	Y N	и и и
0.8	N.E.	N.E.	N	и и	и и и
09	NOT ESTAB	NOT ESTAB	N	Y Y	и и и

SARA 311/312 CATEGORIES FOR THIS PRODUCT: ACUTE= Y, CHRONIC= Y, FLAMMABILITY= Y, PRESSURE= N, REACTIVITY= N

OCCUPATIONAL EXPOSURE LIMITS HAVE BEEN ESTABLISHED FOR THE FOLLOWING MATERIALS:

	WCG1.	D.	0.5.	OBRA
REF	TLV-TWA	TLV-STEL	PEL-TWA	PEL-STEL
01	100 ppm	125 ppm	100 ppm	125 ppm
02	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.
03	100 ppm	150 ppm	100 ppm	150 ppm
04	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.
05. R-	2 mg/m3	NOT ESTAB.	R- 2 mg/m3	NOT ESTAB.
06	400 ppm	500 ppm	400 ppm	500 ppm
07	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.
08 R-	2 MG/M3	NOT ESTAB	2 MG/M3	NOT ESTAB
09	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.	NOT ESTAB.

[C- Ceiling Limit; S- Potential Skin Absorption; R- Respirable Dust] [NOT ESTAB. = NOT ESTABLISHED = NOT APPLICABLE]

PRODUCT STATUS RELATIVE TO THE U.S. EPA TOXIC SUBSTANCES CONTROL ACT

All chemical substances in this product are listed on the U.S. TSCA Inventory or are otherwise exempt from TSCA Inventory reporting requirements.

SECTION 3 - HAZARDS IDENTIFICATION

EFFECTS OF OVEREXPOSURE FROM:

INGESTION: Harmful if swallowed.

EYE CONTACT: This product contains a material which causes irreversible eye damage.

SKIN CONTACT: May cause skin burns. May be absorbed through the skin. Prolonged or repeated contact may cause an allergic skin reaction.

INHALATION: Vapor and/or spray mist harmful if inhaled. Vapor irritates eyes, nose, and throat. Repeated exposure to high vapor concentrations may cause irritation of the respiratory system and permanent brain and nervous system damage.

MSDS # 057950

CHRONIC OVEREXPOSURE: Avoid long-term and repeated contact. Prolonged inhalation of an ingredient(s) in this product may cause edema of the lungs and/or lung damage. This product contains talc. In a lifetime inhalation study female rats exposed to an elevated respirable concentration (9 times the Permissible Exposure Limit) of cosmetic grade talc developed lung cancer. To date, no U.S. regulatory agency has classified talc as a carcinogen based on this data. Ethylbenzene has been reported by NTP to cause cancer in laboratory animals following a chronic (2 year) inhalation exposure. Dose levels of 75, 250 and 750 ppm were used, with evidence of carcinogenicity found in the kidneys of rats and the lung and liver of mice at 750 ppm. The No Observed Effect Level (NOEL) was 75 ppm. The relevance of these findings to humans is uncertain, but appropriate safeguards should be employed to reduce or eliminate inhalation exposure to ethylbenzene. High exposures to xylenes in some animal studies have been reported to cause health effects on the developing embryo and fetus. These effects were often at levels toxic to the mother. The significance of these findings to humans has not been determined.

SIGNS AND SYMPTOMS OF OVEREXPOSURE: Eye watering, headaches, nausea, dizziness, and loss of coordination are indications that solvent levels are too high. Intentional misuse by deliberately concentrating and inhaling the contents can be harmful or fatal. Redness, itching, burning sensation and visual disturbances may indicate excessive eye contact. Dryness, itching, cracking, burning, redness, and swelling are conditions associated with excessive skin contact.

MEDICAL CONDITIONS AGGRAVATED BY EXPOSURE: Not applicable.

SECTION 4 - FIRST AID MEASURES

INGESTION: If swallowed, give one to two eight ounce glasses of water, but do not induce vomiting. Gently wipe out inside mouth to remove any residual material.

EYE CONTACT: In case of eye contact, remove contact lenses, flush eye immediately with a gentle stream of warm water for at least 30 minutes.

SKIN CONTACT: In case of skin contact, flush immediately with plenty of water for at least 15 minutes followed by washing with soap and water.

INHALATION: If affected by inhalation of vapor or spray mist, remove to fresh air. Apply artificial respiration and other support measures as required.

OTHER: If ingestion, any type of overexposure or symptoms of overexposure occur during or following the use of this product, contact a poison control center, emergency room or physician immediately; have Material Safety Data Sheet information available.

SECTION 5 - FIRE FIGHTING MEASURES

FLASHPOINT: 62 Degrees F (17 Degrees C) (PENSKY-MARTENS CLOSED CUP)

FLAMMABLE LIMITS: Lower explosion limit (LEL): 1.8

Upper explosion limit (UEL): Not available

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EXTINGUISHING MEDIA: Use National Fire Protection Association (NFPA) Class B extinguishers (carbon dioxide, dry chemical, or universal aqueous film forming

MSDS # 057950

foam) designed to extinguish NFPA Class IB flammable liquid fires.

UNUSUAL FIRE AND EXPLOSION HAZARDS: Keep this product away from heat, sparks, flame, and other sources of ignition (i.e., pilot lights, electric motors, static electricity). Invisible vapors can travel to a source of ignition and flash back. Do not smoke while using this product. Keep containers tightly closed when not in use. Closed containers may explode when overheated. Do not apply to hot surfaces. Toxic gases may form when this product comes in contact with extreme heat.

SPECIAL FIRE FIGHTING PROCEDURES: Water spray may be ineffective. Water spray may be used to cool closed containers to prevent pressure build-up and possible autoignition or explosion when exposed to extreme heat. If water is used, fog nozzles are preferable. Fire-fighters should wear self-contained breathing apparatus and full protective clothing.

SECTION 6 - ACCIDENTAL RELEASE MEASURES

STEPS TO BE TAKEN IN CASE MATERIAL IS RELEASED OR SPILLED: Provide maximum ventilation. Only personnel equipped with proper respiratory, skin, and eye protection should be permitted in the area. Remove all sources of ignition. Take up spilled material with sand, vermiculite, or other noncombustible absorbent material and place in clean, empty containers for disposal. Only the spilled material and the absorbant should be placed in this container.

WASTE DISPOSAL METHOD: Waste material must be disposed of in accordance with federal, state, provincial, and local environmental control regulations. Empty containers should be recycled or disposed of through an approved waste management facility.

SECTION 7 - HANDLING AND STORAGE

HANDLING AND STORAGE PRECAUTIONS: Do not store above 120 degrees F. (48 degrees C.). Store large quantities in buildings designed and protected for storage of NFPA Class IB flammable liquids.

OTHER PRECAUTIONS: Vapors may collect in low areas. If this material is part of a multiple component system, read the Material Safety Data Sheet(s) for the other component or components before blending as the resulting mixture may have the hazards of all of its parts. Containers should be grounded when pouring. Avoid free fall of liquids in excess of a few inches.

SECTION 8 - EXPOSURE CONTROLS AND PERSONAL PROTECTION

PERSONAL PROTECTIVE EQUIPMENT FOR:

EYE PROTECTION: Wear chemical-type splash goggles or full face shield when possibility exists for eye contact due to splashing or spraying liquid, airborne particles, or vapors.

SKIN PROTECTION: Wear protective clothing to prevent skin contact. Apron and gloves should be constructed of: butyl rubber. No specific permeation/degradation testing have been done on protective clothing for this product. Recommendations for skin protection are based on infrequent contact with this product. For frequent contact or total immersion, contact a manufacturer of protective clothing for appropriate chemical impervious equipment.

MSDS # 057950

RESPIRATORY PROTECTION: Overexposure to vapors may be prevented by ensuring proper ventilation controls, vapor exhaust or fresh air entry. A NIOSH- approved air purifying respirator with the appropriate chemical cartridges or a positive-pressure, air-supplied respirator may also reduce exposure. Read the respirator manufacturer's instructions and literature carefully to determine the type of airborne contaminants against which the respirator is effective, its limitations, and how it is to be properly fitted and used.

OTHER EQUIPMENT: Clean contaminated clothing and shoes.

VENTILATION REQUIREMENTS: Provide general dilution or local exhaust ventilation in volume and pattern to keep the concentration of ingredients listed in Section 2 below the lowest suggested exposure limits, the LEL below the stated limit, and to remove decomposition products during welding or flame cutting.

SECTION 9 - PHYSICAL AND CHEMICAL PROPERTIES

[FORMULA VALUES, NOT SALES SPECIFICATIONS]

BOILING RANGE: 180- 400Degrees F

SOLUBILITY IN WATER: 15.8 %

VAPOR PRESSURE: 19.7 mmHg

WEIGHT/GALLON (LBS): 8.82 (U.S.)

VAPOR DENSITY: Heavier than air

pH: Not applicable

1 - 1 - 1 - 1 - 1

% VOLATILE/VOLUME: 37.830

% SOLIDS BY WEIGHT: 69.25

SPECIFIC GRAVITY: 1.058

EVAPORATION RATE (BuOAc=100): 147

ODOR/APPEARANCE: Viscous liquid with an odor characteristic of the solvents listed in Section 2.

SECTION 10 - STABILITY AND REACTIVITY

This product is normally stable and will not undergo hazardous reactions.

INCOMPATIBILITY (MATERIALS AND CONDITIONS TO AVOID): Avoid contact with strong alkalies, strong mineral acids, or strong oxidizing agents.

HAZARDOUS DECOMPOSITION PRODUCTS: May produce the following hazardous decomposition products when exposed to extreme heat: carbon monoxide; carbon dioxide; lower molecular weight polymer fractions; oxides of nitrogen; Extreme heat includes, but is not limited to, flame cutting, brazing, and welding.

Hazardous Materials Identification System (HMIS) and National Fire Protection Association (NFPA) Ratings:

MSDS # 057950

HMIS Rating	T	NFPA Rating		
HEALTH	3*	HEALTH	3	
FLAMMABILITY	3	FLAMMABILITY	3	
REACTIVITY	0	INSTABILITY	0	

Rating System: 0=Minimal, 1=Slight, 2=Moderate, 3=Serious, 4=Severe, *=Chronic Effects.

Safe handling of this product requires that all of the information on the MSDS be evaluated for specific work environments and conditions of use.

THIS IS THE END OF THE MSDS FOR: 95-2402 (00125842.00195-2402

Manufactured and Supplied by:

PPG INDUSTRIES, EAST POINT

1377 OAKLEIGH DRIVE

EAST POINT, GA 30344

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APPENDIX C ANALYSIS OF PAINT FROM CRANE USED TO SUPPORT MONORAIL, CONCRETE SUPPORT DESIGNATION

From:

Numatec Hanford Corporation

8C530-FAST-97-074

Phone:

373-4771 S3-90 August 5, 1997

Date:

Subject: ANALYTICAL REPORT FOR K-BASIN CRANE REMOVAL - FD1-7021, REVISION 1

To:

R. M. Jochen

X3-67

J. L. Weamer

X3-85

cc: D. J. Smith

S3-90-5-40

FAST File

Reference:

Internal Memo, L. L. Lockrem to R. M. Jochen, "Analytical

Report for K-Basin Crane Removal - FD1-7021, 8C530-FAST-97-

065, dated July 16, 1997.

Please remove and destroy page 6 of the above referenced report and replace with the attached page. The attachment includes additional metals data for samples FD1-7021-01, FD1-7021-02, FD1-7021-04 and FD1-7021-11.

If you have any questions regarding analysis, please contact either Mr. Don Smith at 373-2482 or Ms. Joy Smith at 373-9171.

L. L. Lockrem, Manager Special Analytical Support

sir

Attachment

NHC

Numatec

Hanford Corporation

Internal Memo

An SGN/Cogema, Inc. Company

From:

Special Analytical Support

8C530-FAST-97-065

Phone:

373-4771 S3-90

Date:

July 16, 1997

Subject: ANALYTICAL REPORT FOR K-BASIN CRANE REMOVAL - FD1-7021

To:

R. M. Jochen

X3-67

J. L. Weamer

X3-85

cc: D. J. Smith

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Project File

Attached is the analytical report in support of this project.

If you have any questions regarding analysis, please contact either Mr. Don Smith at 373-2482 or Ms. Joy Smith at 373-9171.

L. L. Lockrem, Manager Special Analytical Support

sir

Attachment

Attachment

ANALYTICAL REPORT

for

FAST PROJECT FD1-7021 K-Basin Crane Removal

Consisting of 17 pages

ANALYTICAL REPORT

for

FAST PROJECT FD1-7021 K-Basin Crane Removal

prepared for

Duke Engineering & Services Hanford, Inc. P.O. Box 350 Richland, Washington 99352

July 15, 1997

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Project Sampling and Analysis Case Narrative

INTRODUCTION

On May 13, 1997, Field Assessment Services Team (FAST) personnel collected paint, oil and grease samples from various locations on a crane that was being removed from the K-Basin area. The analytical data generated was in support of designation and disposal activities. The samples collected were transported with chain of custody to the 622R facility for analysis.

ANALYSIS REQUESTED

Fast Sample ID	Customer ID	Date Sampled	Analysis Requested
FD1-7021-01	 SE Bridgerail	5/13/97	ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se), Total Alpha/Beta
FD1-7021-02	Trolley Body	5/13/97	ICP Metals (Ag_As_Ba_Cd_Cr,Pb,Se), Total Alpha/Beta
FD1-7021-03	Flex Coupling	5/13/97	ICP Metals (Ag_As_Ba_Cd_Cr_Pb,Se), PCBs
) FD1-7021-04	NE Bridgerail	5/13/97	ICP Metals (A <u>g</u> ,As,Ba,Cd,Cr,Pb,Se), Total Alpha/Beta
FD1-7021-05	Pillow Block	5/13/97	ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se)
FD1-7021-06	PillowBlock	5/13/97	ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se)
FD1-7021-07	Flex Coupling	5/13/97	ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se), PCBs
FD1-7021-08	Trolley Gearbox	5/13/97	ICP Metals (Ag, As, Ba, Cd, Cr, Pb, Se), Flashpoint, TX, PCBs, Total Alpha/Beta
FD1-7021-09	Pillow Block	5/13/97	ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se), PCBs
FD1-7021-10	Flex Coupling	5/13/97	ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se), PCBs
FD1-7021-11	Gear & Pinion	5/13/97	ICP Metals (Ag.As,Ba,Cd,Cr,Pb,Se), Flashpoint, TX, PCBs, Total Alpha/Beta
FD1-7021-12	N Bridgerail	5/13/97	ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se), Total Alpha/Beta
FD1-7021-13	Drivetrain Gearbox	5/13/97	ICP Metals (Ag, As, Ba, Cd, Cr, Pb, Se), Flashpoint, TX, PCBs, Total Alpha/Beta
FD1-7021-14	Bridge Gearbox	5/13/97	ICP Metals (Ag As Ba, Cd, Cr, Pb, Se), Flashpoint, TX, PCBs, Total Alpha/Beta
FD1-7021-A	Composite of Samples: 03 & 04		ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se), PCBs
FD1-7021-B	Composite of Samples: 03 & 04		ICP Metals (Ag,As,Ba,Cd,Cr,Pb,Se), PCBs
FD1-7021-C	Composite of Samples: 03 & 04		Total Alpha/Beta

ANALYSIS

Samples FD1-7021-03 and -07 were composited to provide sufficient material for analysis and were similar in matrix and origin.

Samples FD1-7021-05, -06, -09 & -10 were composited prior to digestion since all the waste grease would be combined to allow for efficient and cost effective disposal.

The remaining microwave digest of samples FD1-7021-08, -13, & -14 were composited and analyzed for total alpha/beta levels since the waste oil would be combined to allow for efficient and cost effective disposal.

ICP Metals

The samples were microwave digested and analyzed for the presence of RCRA metals by Inductively Coupled Plasma Atomic Emission Spectroscopy (ICP-AES) using procedure WHC-IP-1128, 3.55. This procedure follows SW-846 Method 6010A.

Flashpoint

Flashpoint was determined using a SET-A-FLASH closed cup flashpoint analyzer following ASTM Method [3]3-90 (reference procedure WHC-IP-1128 3.39 Rev. 0).

Polychlorinated Biphenyls (PCBs)

PCB levels were determined at the Waste Sampling Characterization Facility (WSCF) by Gas Chromatography/Mass Spectrometry using procedure LA-523-457. This procedure follows SW846 Method 8081.

Total Alpha/Beta

Samples were analyzed for alpha/beta levels at WSCF by liquid scintillation.

Total Halides (TX)

Samples were analyzed at WSCF by total combustion in accordance with procedure LA-523-457.

QUALITY CONTROL

All quality control information was within specified acceptable limits except for a high lead spike recovery due to an inadequate spiking level relative to the concentration of lead in the sample and heterogeneity of the sample. In addition, the percent recovery for cadmium in the Laboratory Control Sample (LCS) was above the acceptable limits. This high recovery was most likely due to contamination that occured during sample preparation. Therefore, any positive results for cadmium may be biased high due to possible contamination.

REFERENCES

EPA July 1992, Test Methods for Evaluating Solid Waste (SW-846), Third Edition; U.S. Environmental Protection Agency, Washington, D.C.

WHC-IP-1128, Special Analytical Studies Procedure Manual.

DATA SUMMARY

Project FD1-7021 K-Basin and Removal Results

unple ID		FD1-7021-01	FD1-7021-02	FD1-7021-04	FD1-7021-08	FD1-7021-11	FD1-7021-12	FD1-7021-13	FD1-7021-14	FI)1-7021-A	FD1-7021-B	FD1-7021-0
		SouthEast	Trolley	NorthEast	Trolley	Gear &	North	Drivetrain	Bridge	Composite	Composite	Composite
ustomer ID	1	Bridgerail	Body	Bridgerail	Gearbox	Pinion	Bridgerail	Gearbox	Gearbox	À	В	Ċ
ate Sampled		5/13/97	5/13/97	5/13/97	5/13/97	5/13/97	5/13/97	5/13/97	5/13/97	5/22/97	5/22/97	6/25/97
ime Sampled		947	1005	1026	1057	1110	1125	1130	1335			
laterx		Paint	Paint	Paint	Oil	Oil	Paint	Oil	Oil	Grease	Grease	Oil
igestion Method		Microwave	Microwave	Містомаче	Microwave	Microwave	Microwave	Microwave	Microwave	Microwave	Microwave	Microwave
Metals	Ųnits											
g (Silver)	mg/kg	217	68.8	188	.sau	0,48U	169	0.48U	0.97	0.53	0.53	NR
s (Arsenic)	mg/kg	2.71U	1,38U	1.38U	0.77U	0.70U	1.38U	0.69Ư	0.78U	0.77U	0.77U	NR
a (Barium)	mg/kg	73.8	429	1174	65.6	86.4	210	446	1672	130	199	NR
d (Cadmium)	mg/kg	39.0	25.1	23.7	26.9	16.0	42.9	14.8	19.1	25.9	20.5	NR
r (Chromium)	ing/kg	11900	50500	38600	11.4	9.04	6331	0.24	0.84	54.6	25.9	NR
(Lead)	mg/kg	68400	241000	224000	17.4	4.43	10700	3.63	91.7	604	1940	NR
e (Selenium)	mg/kg	9.70U	4.94U	4.940	2.75 U	2.50 U	4.94U	2.48 U	2.78 U	2.76 U	2.75 U	NR
Jeneral Chemistry												
lashpoint	9;	NR	NR	NR	>200	IS	NR	>200	>200	NR	NR	NR
otal Halogens	mg/kg	NR	NR	NR	500 U	IS	NR	IS	500 U	NR	NR	NR
CIJ's O	mg/kg	NR	NR	NR	1.00 U	1.00 U	NR	1.00 U	<1.50 (#1)	<15.0 (#2)	IS	NR
RadibChemistry	Units			·							anderse en	etarestarestares anno 1997. Transportation de la composition de la
otal Alpha	pCi/g	IS	1.8	1.2 ป	NR	NR	5.7 U	NR	NR	NR	NR	2.70 U
otal Beta	pCi/g	IS	12.0	13,0	NR	NR	11.0 U	NR	NR	NR	NR	9.30
omposite #A: Sam	-los 02 (() 22 a) and ()2 ((0.19-)									
omposite #B; Sam												
Composite #C: Sam				diamtions								
omposite #C: Sam	pies -oa,	-13, & -14 (rema	inder of interow.	ave digestions)								
J' Qualifier: The a	nalyle wa	s not detected at	or below the det	ection limit.								
Clb												
OTE#1: FD1-7021	1-14:	The ECD peak	pattern for this sa	imple does not e	xactly match an	Aroclor 1260 pa	item. If the sam	ple were quanti	tated as Aroclo	1260 the resul	lt would	
		be less than 1.5	ppm. Mass spec	etral data did not	reveal any chlor	inated biphenyls	i,					
OTE #2: FD1-702	l-Λ:				xactiy match an			ple were quanti	tated as Aroclo	1260 the resul	t would	
	1	be less than 1.5	ppin. Mass spec	tral data did not	revent any chlor	inated biphenyls						
S: Insufficient Samp	e to Anal	lyse										——————————————————————————————————————

Reviewed by: M

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QC SUMMARY

HNF-8918, Rev. 0 Quality Control Summary - FD1-7021

As (Arsenic) mg/L FD1-7021-B 0.52 8.12 8.00 94.4 Ba (Barium) mg/L FD1-7021-B 3.58 13.3 8.00 117.0 Cd (Cadmium) mg/L FD1-7021-B 0.37 0.59 0.20 93.6 Cr (Chromium) mg/L FD1-7021-B 0.47 1.36 0.80 105.9 Pb (Lead) mg/L FD1-7021-B 34.9 IS 2.0 NA Streelenium) mg/L FD1-7021-B 0 9.22 8.0 115.3 Chatrix Spike (PMS) Spiked Original Spiked Spiking Element Units Sample ID Sample Conc. Sample Conc. Level % Recovery Ag (Silver) mg/L FD1-7021-B 0.01 0.17 0.20 78.9 As (Arsenic) mg/L FD1-7021-B 0.52 8.25 8.00 96.7 Ba (Barium) mg/L FD1-7021-B 0.37 0.47 0.20 51.0 Cd (Cadmium) mg/L FD1-7021-B 0.47 1.38 0.80 114.3 Pb (Lead) mg/L FD1-7021-B 34.9 IS 2.00 NA	CCV2 0.11 3.82 4.04 0.10 0.41 1.04 4.06 SW-846 Limit 75-125%	7-Recovery 106.5 95.6 100.9 103.9 102.5 103.7 101.4
Ray (Arsenic) mg/L 4.0 3.76 94.0 3.94 98.4 Ba (Barium) mg/L 4.0 4.08 102.0 4.13 103.3 Cd (Cadrinium) mg/L 0.1 0.10 102.4 0.11 105.6 Cr (Chromium) mg/L 0.4 0.41 103.6 0.42 105.6 Pb (Lead) mg/L 1.0 1.02 101.9 1.05 104.9 Se (Selenium) mg/L 4.0 4.18 104.4 4.22 105.6 Laboratory Control Sample (LCS) Element Units Actual Value Found % Recovery Ag (Silver) mg/L 8.00 7.43 92.9 As (Arsenic) mg/L 8.00 8.03 100.4 Cd (Cadmium) mg/L 0.20 0.45 232.4 possible contamination Cd (Cadmium) mg/L 0.80 8.01 101.5 Pb (Lead) mg/L 2.00 2.02 101.0 Se (Selenium) mg/L 8.00 8.22 102.8 Matrix Spike (MS) Spiked Original Spiked Spiking Element Units Sample ID Sample Conc. Sample Conc. Level % Recovery Ag (Silver) mg/L FD1-7021-B 0.52 8.12 8.00 94.4 Ba (Barium) mg/L FD1-7021-B 0.52 8.12 8.00 94.4 Ba (Barium) mg/L FD1-7021-B 0.37 0.59 0.20 93.6 Cr (Chromium) mg/L FD1-7021-B 0.47 1.36 0.80 105.9 Pb (Lead) mg/L FD1-7021-B 0.47 1.36 0.80 105.9 Pb (Lead)	3.82 4.04 0.10 0.41 1.04 4.06 SW-846 Limit 75-125%	95.6 100.9 103.9 102.5 103.7
Ba (Barium) mg/L 4.0 4.08 102.0 4.13 103.3 Cd (Cadmium) mg/L 0.1 0.10 102.4 0.11 105.6 Cr (Chromium) mg/L 0.4 0.41 103.6 0.42 105.6 Cr (Chromium) mg/L 1.0 1.02 101.9 1.05 104.9 Se (Selenium) mg/L 4.0 4.18 104.4 4.22 105.6 Laboratory Control Sample (LCS)	4.04 0.10 0.41 1.04 4.06 SW-846 Limit 75-125%	100.9 103.9 102.5 103.7
Cd (Cadmium) mg/L 0.1 0.10 102.4 0.11 105.6	0.10 0.41 1.04 4.06 SW-846 Limit 75-125%	103.9 102.5 103.7
Cr (Chromium) mg/L 0.4 0.41 103.6 0.42 105.6 Pb (Lead) mg/L 1.0 1.02 101.9 1.05 104.9 Sc (Selenium) mg/L 4.0 4.18 104.4 4.22 105.6 Laboratory Control Sample (LCS)	0.41 1.04 4.06 SW-846 Limit 75-125%	102.5 103.7
Po Clead mg/L 1.0 1.02 101.9 1.05 104.9 Se (Selenium) mg/L 4.0 4.18 104.4 4.22 105.6	1.04 4.06 SW-846 Limit 75-125%	103.7
Sc (Selenium) mg/L 4.0 4.18 104.4 4.22 105.6	4.06 SW-846 Limit 75-125%	
Laboratory Control Sample (LCS) Element	SW-846 Limit 75-125%	
Element	Limit 75-125%	
Ag (Silver) mg/L 0.20 0.20 98.9 As (Arsenic) Ba (Barium) mg/L 8.00 8.03 100.4	Limit 75-125%	
As (Arsenic) mg/L 8.00 7.43 92.9	Limit 75-125%	
Ba (Barium) mg/L 8.00 8.03 100.4	Limit 75-125%	
Cd (Cadmium) mg/L 0.20 0.46 232.4* possible contamination Cr (Chromium) mg/L 0.80 0.81 101.5 Pb (Lead) mg/L 2.00 2.02 101.0 Se (Selenium) mg/L 8.00 8.22 102.8 Matrix Spike (MS) Spiked Original Spiked Spiking Belment Units Sample ID Sample Conc. Level % Recovery Ag (Silver) mg/L FD1-7021-B 0.01 0.18 0.20 82.4 As (Arsenic) mg/L FD1-7021-B 0.52 8.12 8.00 94.4 Ba (Barium) mg/L FD1-7021-B 0.37 0.59 0.20 93.6 Cr (Chromium) mg/L FD1-7021-B 0.47 1.36 0.80 105.9 Pb (Lead) mg/L FD1-7021-B 34.9 IS 2.0 NA Spikelenium) mg/L FD1-7021-B 0 9.22 8.0 115.3	Limit 75-125%	
Cr (Chromium) mg/L 0.80 0.81 101.5 Pb (Lead) mg/L 2.00 2.02 101.0 Se (Selenium) mg/L 8.00 8.22 102.8 Matrix Spike (MS) Spiked Original Spiked Spiking Element Units Sample ID Sample Conc. Level % Recovery Ag (Silver) mg/L FD1-7021-B 0.01 0.18 0.20 82.4 As (Arsenic) mg/L FD1-7021-B 0.52 8.12 8.00 94.4 Ba (Barium) mg/L FD1-7021-B 3.58 13.3 8.00 117.0 Cd (Cadmium) mg/L FD1-7021-B 0.37 0.59 0.20 93.6 Cr (Chromium) mg/L FD1-7021-B 0.47 1.36 0.80 105.9 Pb (Lead) mg/L FD1-7021-B 34.9 1S 2.0 NA Selenium) mg/L FD1-7021-B 0 9.22 8.0 115.3 <tr< td=""><td>Limit 75-125%</td><td></td></tr<>	Limit 75-125%	
Pb Lead mg/L 2.00 2.02 101.0	Limit 75-125%	
Sec Selenium mg/L 8.00 8.22 102.8	Limit 75-125%	***************************************
Matrix Spike (MS) Spiked Original Spiked Spiking Element Units Sample ID Sample Conc. Sample Conc. Level % Recovery Ag (Silver) mg/L FD1-7021-B 0.01 0.18 0.20 82.4 As (Arsenic) mg/L FD1-7021-B 0.52 8.12 8.00 94.4 Ba (Barium) mg/L FD1-7021-B 3.58 13.3 8.00 117.0 Cd (Cadmium) mg/L FD1-7021-B 0.37 0.59 0.20 93.6 Cr (Chromium) mg/L FD1-7021-B 0.47 1.36 0.80 105.9 Pb (Lead) mg/L FD1-7021-B 34.9 IS 2.0 NA Scelenium) mg/L FD1-7021-B 0 9.22 8.0 115.3 Element Units Sample ID Sample Conc. Sample Conc. Level % Recovery As (Arsenic) mg/L FD1-7021-B 0.52 8.25 8.00 96.7 <td>Limit 75-125%</td> <td>odčnostoskohorovano i i i i i i i i i i i i i i i i i i i</td>	Limit 75-125%	odčnostoskohorovano i i i i i i i i i i i i i i i i i i i
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Pb (Lead) mg/L FD1-7021-B 34.9 IS 2.0 NA	75-125%	
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Matrix Spike (PMS) Spiked Original Spiked Spiking	75-125%	
Element Units Sample ID Sample Conc. Sample Conc. Level % Recovery Ag (Silver) mg/L FD1-7021-B 0.01 0.17 0.20 78.9 As (Arsenic) mg/L FD1-7021-B 0.52 8.25 8.00 96.7 Ba (Barium) mg/L FD1-7021-B 3.58 13.3 8.00 121.5 Cd (Cadmium) mg/L FD1-7021-B 0.37 0.47 0.20 51.0 Cr (Chromium) mg/L FD1-7021-B 0.47 1.38 0.80 114.3 Pb (Lead) mg/L FD1-7021-B 34.9 IS 2.00 NA	75-125%	
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	75-125%	
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Blanks		
	Prep Blank	
Ag (Silver) mg/L 0.003 0.002 0.00 0.00 0.00	0.06	<u></u>
As (Arsenic) mg/L 0.023 -0.026 0.03 0.02 0.02	2.04	
Ba (Barium) mg/L 0.000 0.002 0.00 0.01 0.00	0.11	*possible
01/01/102	2108	-
Cd (Cadmium) mg/L 0.001 0.004 0.00 0.00 0.00	6.19*	contaminant
Cr (Chromium) mg/L 0.000 0.000 0.05 0.03 0.00	0.02	
Pb (Lead) mg/L 0.021 0.002 0.24 0.16 0.00	-0.48	
Se (Selenium) mg/L -0.012 0.010 -0.02 -0.01 -0.01	0.70	
Duplicates MS & PMS	and the control of the second	
Element Units Sample ID MS PMS % RPD		
Ag (Silver) mg/L FD1-7021-B 0.18 0.17 3.99		
As (Arsenic) mg/L FD1-7021-B 8.12 8.25 1.59		
3enium) mg/L FD1-7021-B , 13.3 13.3 0.00		
(Cadmium) mg/L FD1-7021-B 0.59 0.47 23.0		
Cr (Chromium) mg/L FD1-7021-B 1.36 1.38 1.46		
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Se (Selenium) mg/L FD1-7021-B 9.22 9.57 3.73	-	<u></u>
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NA = Not Applicable	-	

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Printed on: 7/16/97

CHAIN-OF-CUSTODY INFORMATION

NF-8918, Rev. 0

Treful Assessment Services
Special An fical Services
Hanford Te cal Services
Richland, WA. 99352
(509) 373-3798 FAX (509) 373-3193

Chain-of-Cistody/ Sample Analys Request

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FB1-7021-02	5-15	10:05	So	11 Horsen Ord	1		-			1			TF.	المعلامة 11 س م	Bily (Pilin	4)	
FB1-7021-03	5-13	10:15	0	1, 125 AG	- -				-	 			/L-1	: -:	ex civaling	RIII	
1701-7021-04	5-13	10:26		1 plate	-								Br	ide R.	Body (Paint ex coupling, " INE E	المار والم	
FQ1-7-21-05	5-13	14:20	0	1 20ml CG	_			-					3-	F)	<u> </u>	F	
EDI-7-21-06	5-13	10:20	B	1 20 ml CE									1 1-	•	I.u Blick		
FB1-7021 - 07	5-13	10:42		1, 2mL CC									16.0	T) Fle	x Carplan		
FD1-7021 - 08	5-13	10-57	0	1-125 ml LP	5								IR	//x 6	err Bix	(1-0)	
FD1-7021-09	5-13	11:06	O	ŧ;											Hu Blick		
FBI-7021-10	5-13	11:08	0	L 20ml Co										•	lex coupling		
FOI - 7021 - 11	5-13	11:10	0	1,125 AB									(4.	G) G	cur i Pinie		
FBI - 7021 - 12 Special Shipment/1	5-13	11:25	50	1125 AG									B_{E}	dic Bai	eur flinien L Northerd	Post	
Special Shipment/1 Storage Requireme	**	or	-		ı	isibl marl		mple	lla	zar	ls/		— ,-	1		☐ Ra	d Screening
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PICIO" MSSCSSMCHC" SCTVICCS Team

Special Ar , 'ical Services

Hanford To .cal Services

Richland, WA. 99352

(509) 373-3798 FAX (509) 373-3193

Chain-of-Cistody/ Sample Analy Request

SAS II	· ,
Date 5-13.97	1
Date	

Page 2 01 2

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Disposition																		

SUPPLEMENTAL INFORMATION

WSCF (S) ANALYTICAL LABORATORY REPORT

Attention:	
Project Number	

Joy Smith FAST

:FAST 1997

Group #:

97000885

nple#	Client ID	Test Performed		-Matrix	Method	RQ	Result	Units	MDL	Sampled	Received	
70002164	F01-7021-02	Alpha by liquid scintillation		SOLID	LA-508-42		1.8	pCi/g	0.80	06/10/97	06/10/97	
70002164	FD1-7021-02	Alpha error by LC		SOLID	LA-508-42		120	%	0.00	08/10/97	08/10/97	
70002184	F01-7021-02	Beta by liquid scintillation	10.0	SOLID	LA-508-42	i de la companya da l	1.28+01	pCi/g	1.60	06/10/97	06/10/97	
77702164	F01-7021-02	Bola error by LC		solip 👙	LA-508-42		40	ж.	0.00	06/10/97	06/10/97	
J2165	FD1-7021-04	Alpha by liquid scintillation	a season was a season	SOLID	LA-508-421	U	4,0E-01	ρCi/g	1.20	06/10/97	06/10/97	
70002165	FD1-7021-04	Alpha error by LC	a sykyy a digyyddd Y achel a argaillau y	SOLID	LA-508-421		680	%	0,00	08/10/97	08/10/97	
70002165	FD1-7021-04 •	Beta by liquid scintillation	arende kriti uzben ditabile	SOLID	LA-508-421	Mada riyar bisa mi '	1.3E+01	pCi/g	2.20	06/10/97	06/10/97	
70002165	F01-7021-04	Date error by LC		SOLID	LA-508-421	98.9843 S	50	%	0.00	06/10/97	06/10/97	
70002166	FD1-7021-12	Alpha by liquid scintillation		SOLID	LA-508-421	U	5.7	pCi/g	7.60	06/10/97	06/10/97	
70002160	FD1-7021-12	Alpha error by LC		SOLID ////	LA-508-421	Market	320	n	0.00	00/10/97	08/10/97	
770002166	FD 1-7021-12	Beta by liquid scintillation	i santan nela	SOLID	LA-508-421	U	1.0E + 01	ρCi/g	14.00	06/10/97	06/10/97	H
70002166 -70002166 -70002166	FD1-7021-12	Bata arror by LC		20TIO	(A-500-421		.330	∳% J\$``\\$	· 0.00	06/10/97	06/10/97	HNF-8918, Rev. 0

MDL=Minimum Detection Limit

RQ=Result Qualifier

B - The analyte was detected in the associated method blank.

E - Compound concentration exceeded calibration range.

N - Identification is based on a mass spectral library search.

D - Compound concentration resulted from a dilution.

J - Estimated value.

Z - Soe Commonts.

U - The analyte was not detected at or below detection limit.

W004.revI

Field Assessment Services Team

^{* -} Indicates results that have NOT been validated.

HNF-8918, Rev. 0

ANALYTICAL LABORATORY REPORT

Attention: **Project Number** Joy Smith FAST

:FAST 1997

Group #:

97000828

unple#	Client ID	Test Performed	Matrix	Method	RQ	Result	Units	MDL	Sampled	Received
970001903	FD1- (021-08	PCB's in ails	OTHER	EPA 8081	U	U	mg/kg	1,00	05/28/97	05/28/97
970001903	FD1-7021-08	Total Helides	OTHER 🥢	EPA 9076	· <	500	იე/ე	50.00	05/28/97	05/28/97
970001904	FD1-7021-A	PCO's in oits	OTHER	EPA BOB1	z <	15	mg/kg	1.00	05/28/97	05/28/97
101905	FD1-7021-03	PCB's in oils	OTHER	EPA 8081	Z <	(C 15	mg/kg	1.00	05/28/97	05/28/97
01906ء ۽	FD1-7021-B	PCD's in oils	OTHER	EPA 8081		2	mg/kg	1.00	05/28/97	05/28/97
970001907	FD1-7021-14	PCB's in oils	OTHER	EPA 8081	. Z 🦠 <	1.5	mg/kg	1.00	05/28/97	05/28/97
970001907	FD1-7021-14 •	Total Halides	OTHER	EPA 9076	<		սց/ց	50.00	05/28/97	05/28/97
970001508	FD1-7021-11	PCOs for swipes and smears	OTHER	EPA 8081	ំប	0	nīg/kg	1.00	05/28/97	05/28/97
970001909	FD1-7021-13	PCBs for swipes and smears	OTHER		U	0	mg/kg	1.00	05/28/97	05/28/97

MDL=Minimum Detection Limit

RQ = Result Qualifier

B - The analyte was detected in the associated method blank.

E - Compound concentration exceeded calibration range.

N - Identification is based on a mass spectral library search.

D - Compound concentration resulted from a dilution.

J - Estimated value.

Z - See Comments,

U - The analyte was not detected at or below detection limit.

* - Indicates results that have NOT been validated.

W004.rev1

Field Assessment Services Team

2

HNF-8918, Rev. 0

WSC ANALYTICAL COMMENT REPORT

Attention: Project Number Joy Smith FAST

Group #:

97000828

mple#	Client ID	Lab Aı	rea Test	Comment
970001903	FD1-7021-08	VALTEST	Total Halides	TX: Samples were mixed prior to analysis. Each
	4			sample was analyzed four times. W970001903: Average -
-				172.9 ug/g; RSD + 37.3%. W970001907: Average + 113.7 ug/g;
				RSD - 51,8%. Although a large variation was seen, results
				are wall below the regulatory limit of 1000 ug/g.
970001904	FD1-7021-A	ATESTDAT	ra PCB's in oils	The ECD peak pattern for this sample does not exectly match
		•		an Aroclor 1254 pattern. If the sample were quantitated as
		William Committee		Aroclor 1254 the result would be less than 15 ppm.
970001905	FD1-7021-03	TESTDAT		The ECD peak pattern for this sample does not exactly match
	$\mathcal{I} = \mathbb{S}(1, 2, 2, 2)$			en Aroclor 1254 pattern. If the sample were quantitated as
				Aractor 1254 the result would be less than 15 ppin. Mass
Ċ				spectral data for this sample did not reveal any chlorinated
C-20				biphenyls.
970001908	FD1-7021-B	TESTDAT	A PCB's in oils	The ECD peak pattern for this sample does not exactly match
				an Aroclor 1248 patturn. If the sample were quantitated as
• 1.				PCB the result would be less than 2 ppm. Matrix spike(1248)
•				recovery was 95%.
970001907	FD1-7021-14	TESTDAT	A PCB's in oils	* The ECD peak pattern for this sample does not exactly metch
				an Aroclor 1260 pattern. If the sample were quantitated as
				Aroclor 1260 the result would be less than 1.5 ppm.
			,	

Lab Areas:

VALGROUP - Group Validation LOGSAMP - Login for Sample

VALTEST - Test Validation LOGTEST - Login for Tests

TESTDATA - Test Data Entry

Braden, Janis K

From:

Watson, David J (Dave)

Sent:

Friday, November 02, 2001 11:47 AM

To:

Subject:

Braden, Janis K FW: Request to Reproduce a WSCF Analytical Report

----Original Message---

From: Sent:

Fitzgerald, Scot L

To:

Friday, November 02, 2001 11:03 AM Watson, David J (Dave)

Cc: Subject: Powell, Katherine L; Dale, Troy F; Trechter, John E Jr. FW: Request to Reproduce a WSCF Analytical Report

Dave,

I approve the reproduction of this report for the purposes stated below. If you need a form of approval other than this email please let me know.

Scot Fitzgerald Analytical Manager WSCF Laboratory 373-7495

----Original Message----

From:

Powell, Katherine L

Sent:

Friday, November 02, 2001 7:46 AM

To:

Fitzgerald, Scot L

Cc:

Trechter, John E Jr.; Dale, Troy F

Subject:

FW: Request to Reproduce a WSCF Analytical Report

----Original Message-----

From:

Watson, David J (Dave)

Sent:

Friday, November 02, 2001 7:38 AM

To: Cc: Powell, Katherine L Braden, Janis K

Subject:

Request to Reproduce a WSCF Analytical Report

We are using WSCF "Analytical Report for FAST Project FD1-7021 K-Basin Crane Removal" in Revision 0 ot the Sampling and Analysis Plan for Structures External to the 100K Storage Basins", HNF-8918. This report is included in its entirety as an appendix and is referred to in the text of the SAP as providing historical information on past characterizations.

Within that analytical report there is a sheet titled, "WSCF ANALYTICAL COMMENT REPORT" for Group 97000828 that has on the bottom of the page the following statement: "This report may not be reproduced, except in its entirety without the written approval of the WSCF Laboratory."

As the report is being included in its entirety in the above HNF docuement, copy of which was provided to you in a October 19 meeting, our document clearence staff need to have your approval to reproduce it for release. Can you please provide.

Thanks

Dave Watson 373-3250

HNF-8918, Rev. 0

WSCF () ANALYTICAL LABORATORY REPORT

Attention:

Joy Smith

Group #:

97000989

nple#	Client ID	Test Performed	Matrix	Method RQ	Result	Units	MDL	Sampled	Received
/0002327	FD1-7021-C	Gross Deta	OTHER	LA-508-410	9.3	pCi/g	2.70	06/25/97	06/25/97
70002327	FD1-7021-C	Gross Deta % Method Error	ОТНЕП	LA-500-410		%	0.00	00/25/97	. 00/25/97
	FD1-7021-C	Total Alpha	ОТНЕЛ	LA-508-410 U	-3.9E-01	pCi/g	2.70	06/25/97	06/25/97
102327	F01-7021-C	Total Alpha % Method Error	лэнто	LA-508-410	350	%	0.00	06/25/97	06/25/97

C-2

MDL=Minimum Detection Limit

RQ=Result Qualifier

B - The analyte was detected in the associated method blank.

E - Compound concentration exceeded calibration range.

N - Identification is based on a mass spectral library search.

D - Compound concentration resulted from a dilution.

J - Estimated value.

Z - Soe Comments.

U - The analyte was not detected at or below detection limit.

W004.revI

Field Assessment Services Team

^{* -} Indicates results that have NOT been validated.

APPENDIX D RADIOLOGICAL SURVEY K000511, "INVESTIGATIVE SURVEY OF OLD RAD PAD"

Originally it was reported by K Basin staff that all of the hot spots currently identified by survey on the Filter Wash Pad would not be part of this removal action. Subsequent meetings determined that some of the spots would be left behind but others would be removed. The survey data were evaluated to assess whether disposal of higher contaminated areas would allow the use of the current curie-to-weight ratios. The evaluation assessed whether current ratios would underestimate to the Cs-137 content.

The discussion below illustrates that even if all of the highest contaminated concrete squares scheduled to be removed are placed in one waste box, the weight to curie conversion factor currently used by K East Basin for above water waste will be conservative. Using the factor will not underestimate the Cs-137 in the waste from this action.

The figure attached to the survey provides 3'x3' square grids that are uniquely numbered. The numbers in this paragraph are grid numbers. The area of the Filter Wash Pad that is being removed is divided from the area being left by a north south line between squares 315 and 316 on the north and squares 15 and 16 on the south on the map in the attached survey report K000511. Squares 66,67,68,69,91 are in the area to be removed and they are all >1,000,000 dpm/100cm2. Three other squares (40,65,90) are immediately adjacent to the line and are all listed as >1,000,000 dpm per 100 cm2 but are not scheduled for removal.

It was assumed that all five of the highest contaminated squares of concrete from the area to be removed were placed in a 4x4x8 ft wood box. The 4x4x8 ft wooden box has a dose to Cs-137 curie conversion factor published in WHC documents (WHC 1996a, 1996b). If the five squares of concrete are placed in the box, the maximum dose rate on the outside of the box (ignoring shielding by the box or self shielding by the concrete) is assumed to be a simple addition of the field dose rate survey (≥30 cm from slab) with the dose rate meter window closed.

The closed window reading at 30 cm was selected because it is the dose rate of gamma emitters on the concrete. All gamma emitters are assumed to be Cs-137 for this exercise. The data from the survey sheet for the five squares that are in the area to be removed (squares 66, 67, 68, 69, 91) indicate dose rates of 2.7,0.7, <0.5, <0.5 mR/hr. If the results are summed the in-situ survey dose on the five portions of the slabs equal 4.4 mR/hr.

After placing the contaminated concrete in a waste box, the average dose rate on the waste box could be measured. A rough estimate of how the dose rates from the contaminated concrete would be reduced by the shielding of the other concrete and the wood box was made. It was estimated that the dose rate outside of a wood box full of rubble would be that less than 10% of the dose rate measured when the survey reading is taken 30 cm away from the in-situ concrete without shielding. Applying a 10% factor for shielding and geometry, one could estimate an average measured dose rate outside of the hypothetical box of waste (averaged over all sides of the box) of 0.44 mR/hr (4.4E-4 R/hr). The dose rate conversion for the wood box is 13.8 Ci Cs-137 per R/hr. Thus, if the five contaminated concrete squares were placed in a 4x4x8 wooden box, and an average reading of 0.44 mR/hr was obtained, it is estimated that the five contaminated squares contain about 6.1E-3 Ci of Cs-137.

Using the current K East weight to curie methodology, the concentration of Cs-137 in the waste that is assumed for all of K East above water waste is 1.12E-5 Ci/kg. According to the current K Basin SAP in use (HNF 2001), an ERDF roll-off box can be loaded with 7,980 kg. Applying the current factor to this maximum load, a Cs-137 amount of approximately 9E-2 Ci of Cs-137 would be within the profile. If one estimates the number of curies in the five contaminated sections (6.1E-3 Ci) and divides by the maximum amount that can be placed in the ERDF box, the Cs-137 content of the waste container is nearly a factor of 15 below the maximum allowed in the current profile. Even if the boxes were sent to ERDF only half full, the waste would not exceed the profile.

The calculations and assumptions presented above are conservative in that it is not expected that all of the highest contaminated portions of the pad will end up in one ERDF roll-off box. The highly contaminated section of the pad is less than 4% of the total of the surface area of the pad. It is also known that the dose to Cs-137 curie conversion factors (WHC 1999a, WHC 1999b) are conservative, so it is likely that the 10% factor that was used to convert the dose measured in the survey to a dose that would be measured if the waste were put in a 4x4x8 wood box, would actually be less. That would translate into a lower number of Curies of Cs-137 being placed in the ERDF roll-off box and provide an even larger safety margin.

		Мар	o/Sketch			SNF PROJECT RADIOLOGICAL SURVEY REPORT Pag	re 1 of <u>]</u>
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α	Рт	α	L	βγ	α	βγ	a		βγ		mrem	mrem/hr	/hr	mren		μ 		/414	una	V/111	,,,,,	cm			
NA P	71,000,000	NA	↓_	4/000	NA	70,000	/A	1	20,000	9	43	88.7 N.4	١	NA	5/	JA.	6	9	1.3	NA	15	S/F	·	Grid 91	1
	10	_	<u> </u>	< 1000		3500		_	3500	A -	NA	NA		•	NA	A	N	<u> </u>	N	JΑ	<u>Lr</u>	NA	•	Grid 92	2
<u>'</u>	60,000		\downarrow	<1000		4000	$\downarrow \downarrow$	上	3000			·	\bot			Ц.					<u> </u>			Grid 93	3
1	4D		_	< 1000		2000		_	2000						\perp									Grid 94	9
-	30,000		$oldsymbol{ol}}}}}}}}}}}}}}$	<1000		2000	\coprod		1000		,												<u>.</u>	Grid 95	5
1	4D.	_	L	≺ /000		1000		_	1000														,	Grid 96	6
	< D		1_	<1000		600	\coprod		600					,									<u> </u>	Grid 97.	7
 	< D			< 1000		550		L	550															Grid 98	3
1	<d td="" <=""><td>\perp</td><td>_</td><td><10∞</td><td></td><td>700</td><td></td><td></td><td>700</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td><u> </u></td><td>Grid 99</td><td>2</td></d>	\perp	_	<10∞		700			700														<u> </u>	Grid 99	2
	< 0	\perp	1	<1000		700	Ц	_	700												<u> </u>		0	Grid 100	2
<u> </u>	420,000	_	\perp	<1000		20,000		_	3000											\perp	<u> </u>		<u> </u>	Grid 101	<u> </u>
	₹D,	_	igspace	∠1000		1500		Ļ	1500						<u>.</u>						<u> </u>		2	Grid 102	2
	15,000	1.	_	<1000		2000	\coprod	$oxed{\bot}$	1000												<u> </u>		3	Grid 103	3
\	50.000		$oldsymbol{\perp}$	< 1000		2500		_	1000						_								<u> </u>	Grid 104	4
의	700.000		-	< 1000		4500		1	1000				\mathcal{I}	2ζ	, -	4		-		\perp	<u></u>		<u>S</u>	Grid 105	<u> </u>
	30.000	<u> </u>	4	<1000		2000	$\perp \mid$	 	1000				: ;	7			7 (-				Grid 10b	2
	15,000	\perp	-	000</td <td> </td> <td>1500</td> <td>- - </td> <td>╀-</td> <td>1000</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>1</td> <td>(1)</td> <td></td> <td></td> <td>1-1</td> <td></td> <td>_</td> <td>·· · · · · · · · · · · · · · · · · · ·</td> <td>erid 107</td> <td><u>'</u></td>	 	1500	- -	╀-	1000							1	(1)			1-1		_	·· · · · · · · · · · · · · · · · · · ·	erid 107	<u>'</u>
	15,000		-	< 1000 ⋅		1500	\coprod	╁	1000			<u>-</u>			_ _	_ -		•						Gn d 108	8
	15,000	-	\vdash	41000	<u> </u>	1500	\sqcup	<u> </u>	1000				·			_				-				<u> srid 109</u>	9
	324,000	-	+	<1000		6000	\perp	4	600						_ _	_ _						 		Grid 110	2
_ I	378,000			<1000	 	7000	-	-	700	\square					_ _		-			\sqcup		_		Grid 111	_
	378,000	-		<1000		7000	$\perp \perp$	<u> </u>	700				_		_ _	_	-			-		 ·		Grid 112	
	90,000			< 1000		3000	- -	1-	1500	\vdash						4						_		<u>Grid 113</u>	. 1
	10,000		_	< 1000		1500	Ц	1	1000				\dashv			_				1-1	<u> </u>			Grid 114	_
00	45,	NΑ		<1000	NA	2500	ΙĄ		1500	A	ν.	NA	+	N	NA	Ä	h	Ą	N	Ā	N	Au		Grid 115	

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SN	F PROJECT RADIOLOG	GICAL	SUF	RVEY	REF	וסי	?T -	Co	ntir	ıua	tion	She	et	- Data	Surve		nno. 000511			Page	7_ of	16
ĺ	·			N	ote 1 F	OSI = Fid	E RATI eld (≥ 3	O cm	ASU:) C	REMI = Co	ENTS ntact	(<u><</u> 1 cm	1)				CONT	FAMINAT	TION MEASU	REMENT	s	
la.	Description	, No		WO mR/hi	VA ID F		CF _β	С	F _y	D٥		Shalk Dos	e	Deep Dose	ct	kg om	Gross	Direct	Remo dpm/10	vable 00 cm ²	To dpm/1(ial 00 cm²
_		c	m		_			_		mrei	TIZOZ	mrem	/nr	mrem/hr	βγ	α	ργ	α	βγ	α	βγ	α
6	Grid 116	<u> </u>	Ą	NA	<u> </u>	<u> </u>	NA	1	JA	<u>.</u> N	A	N	4	NA	3000	٩٧٨	4000	NA	< 1000	NA	30.000	<u> </u>
17	Grid 117		1	-		↓_		┿-	\perp				_		2000		3000		<1000		30,000	
8	Grid 118	<u> </u>				\perp		-	$\bot\!\!\!\!\!\bot$						2.000		2000		< 1000	-	< D	ļ
9	<u>6rid 119</u>	l			<u> </u>	╀			\sqcup		$\downarrow \downarrow \downarrow$				1000		1000	<u> </u>	< 1000		< D_	
20	<u>Grid</u> 120		\sqcup		_	<u> </u>	\sqcup	1			 		_		800.		800	-	<1000	<u> </u>	<d< td=""><td></td></d<>	
4	<u>Grid 121</u>				 	L		╀					_		600		600		<1000		< D	<u> </u>
22	<u>Grid</u> 122 ·			_	 	igspace		<u> </u>	_		\sqcup		_		-500		500		<1000		< D.	
3	Grid 123			<u>_: </u>	↓_	_		1					_		400		400		<1000		<0	
4	Grid 124					ļ_							_	_	500		500		<1000		<1>	·
35	Grid 125				┷	_		_					_		500		500		21000		<d< td=""><td></td></d<>	
6	Grid 126	_											_		550		550	<u> </u>	< 1000		< 17	
1	Grid 127 :				<u> </u>				Ц.						550		550		< 1000		< <u>D</u>	
8	Grid 128							$oldsymbol{ol}}}}}}}}}}}}}}}}}}$							600		600		41000		<d< td=""><td></td></d<>	
7	Grid 129														700		700		<1000		< D	
0	Grid 130				<u> </u>						7	7			650		650		<1000		42	
1	Grid BI						\ \)				T		700		800		41000		6,000	
_	Grid 132					X	<u>}</u>	\``\	' '	3					900		20,000		41000		646,000	
3	Grid 133														900		3000		4/000		52,000	
34	Grid 134											•			500		500		<1000		< D	
5	Grid. 135														500		500		4/006		۷D	
6	Grid 136			. .				·							500	\top	560		41000	$\neg \uparrow \neg$.4D	
	Grid 137	•				П									600		700		∠1000		6,000	
	Grid 138														1000		1000		000</td <td></td> <td>< D</td> <td>•</td>		< D	•
	6rid 139									• •					1000	7	1000		<1000		< D	
	Grid 140	N	$\langle \top$	NA	7		NA	N	A	N	A	NA	7	MA	1000	ΝA	1000	NA	000</td <td>νA</td> <td>40</td> <td>NA</td>	νA	40	NA

SN	F PROJ	IECT RAI	DIOLOGICA	AL S	UĘ	₹VE\	/ R								t -	Data	Surve		it No. 00511			ŀ	Page	2 of	jp .
							lote 1	009 F = F	E RA leid (≥	TE N 30	/EAS	UREM C = Co	ENTS ontacl	s (≤1 cm)					, (* CON	(amina	TION MEAS	JREM	IENT:	5	
No.		Description	on '	Dis Not on	e i l	WO mR/h		WC mR/hr	CF	a	CFy	0	utron ose	Shallov Dose	1	Deep Dose	ct B	kg om	Gross	s Direct n/PA	Ren dpm/	novab 100 ca	le m2	To dpm/10	etal 00 cm²
_			<u> </u>				_			_		mre	m/hr	mrem/h		nrem/hr	βγ	a	Рγ	α	βγ		α	βγ	CL 673100
141	Grid	141		NA	+	NΑ	4	NA	N	A	NA	Λ	/A	NA	_	NA	1500	NA	1500	NA	< 1000	1 ^	/ <u>A</u>	<0	< D N
142	Gnid	142		-		-	_					<u> </u>			\perp	_	1000		1000		- 1000	1	\perp	< D	< 4
143	Gr. d	143		-				-	<u> </u>			<u> </u>					750		750	 	<1000	<u> </u>		<:D	< 1
144	Gri d	144	· -		_		\perp		<u> </u>			<u> </u>			\perp		450		450		<1000	<u> </u>		< D	<u> < ∤</u>
45	Grid	145						_	-						_		400		400		< 1000	<u>.</u>		<d< td=""><td><></td></d<>	<>
46	Grid .	1460	·		_		4	\perp		_							350		350	<u> </u>	00</td <td>2</td> <td></td> <td>< D</td> <td>4></td>	2		< D	4>
47	Grid	147			4		_	ŀ		\downarrow					_		350		350		≺/00¢		1	<d< td=""><td>< D</td></d<>	< D
48	Grid	148					_ _	<u> </u>		_				<u> </u>	\perp		400		400		<1000	1.		<d< td=""><td>4P</td></d<>	4P
49	Grid	149								_					\perp		400		400		< 1000	1	1	40	140
150	Grid	150					1			1				<u>L</u>			400		400		1000</td <td>1_</td> <td><u> </u></td> <td>< D</td> <td>4 D</td>	1_	<u> </u>	< D	4 D
151	Grid	151			_	-				-	\perp						600		600		<1000	<u> </u>		<0	* D
52	Grid	/52			_	<u>· </u>	_					<u> </u>					400		400		<1000			< D	KD
153	Grid.	<u> 153</u>	·····		1		\perp			\Rightarrow	TE	27.	\leq				450		450		< 100	2	<u> </u>	4D	\$D
154	Grid	154	·		_		\perp	111			Щ		11				450		450		< 1000		_	∠ D	KD.
22	Grid.	155			_		_ _	1/2	2	¥							450		450		<100	2		< D	£ 18.
56	Grid	156	;		_										┸	_	500		500		< 1000			< D	3€.F.
57	Grid	157	·	_	_					_	1						600		2000		< 1001			84,000	ļ
158	<u>Grid</u>	158			_												700		700		₹/000			∠D	<u> </u>
59	Grid	159		ΝΆ		NA		νА	NA	_	NA			NÀ		ΝA	750		750		<1000			4D	<u> </u>
60	Grid	160	>3"disc	آ را		8 N		·5 20.5	6/2					48 NA	1	0.5	600		50,000		< 1000			7/,000,000	
61	Grid	161		NF	+	NΑ		VΑ	NA	L	NA			NA		NA	600		600		<1000			4 D	
62	Grid	162		NΑ		NA		JA	NA		NΑ			NΑ		NA	650		550		<1000			4D	
63	<u>Grid</u>	163	73"disc		<u>- 1</u>	3.b./		20.5	6N					21.6 NA	40	- Z 0.7 I	650		35,000		<1000			71,080,000	
,4	grid	164	73"disc	رعب	EL	8 N	1	20.5	60		-			47.3 N	A 3	5/0.5	850		50,000		<1000			7/,800,000	
65	Grid	165	• •	NA	+	NA		MA	NV	۲] ٔ	NΑ	7	A	NΑ		NA	900	NA	1000	NA	41000		VΑ	6,000	

SN	F PROJECT RA	DIOLOGICA	L SUF	RVE	ΥR	EP	OR	Τ-	Co	nti	nua	tion	She	et	- Da	ita	Survey		nt No. (00051	İ		Pag	e <u>9</u> of	16
		,			Note	_ 1 F ±	OSE Field	RAT d (≥	E M 30 c	EASU	JREMI C= Co	ENTS ntact	(<u><</u> 1 cm	1)		,			CONT	AMINATIO	ON MEASÚR	EMENT	rs 	
No.	Descrip	lion	Dist.	w		W		CF		CF _y	Neu	tron	Shall Dos		De Do	ep se	ch Bl			Direct n/PA	Remo dpm/10	vable 0 cm2	Tot dpm/10	al 0 cm ²
			cm	mR/	nr)	mR/	nr	-	3	- γ	mre	m/hr	mren	ı/hr	mrei	m/hr	βγ	α	рү	α	βγ	α	βγ	α
66	Grid 166		ΝA	NA	1	NA		ΝĀ	Ц	NA	N	A_	NA	<u>\</u>	_ ∧	<u>A</u>	700	MA	700	NA	<1000	NA	12D	<u> </u>
67	Grid 167						_		_ _			<u> </u>					500	_	500		<1000		< D	
168	Grid 168										<u> </u>						500	_ _	500		000</td <td>· </td> <td>< D</td> <td>- -</td>	·	< D	- -
69	Grid 169												<u> </u>				600		600		<1000		40	\rightarrow
70	Grid 170								\perp								500	_ _	500		< /000	-	< 1)	
71	Orid 171								ŀ	_							450	_ _	450		<1000		4D	\dashv
72	Grid 172								_		,				ده د ژه		350		350		₹1000		40	
13	Grid 173.		!							\perp			<u> </u>				300	_ _	300		<1000		≺ D	$-\!\!\!+$
174	Grid 174											L_					300	_	300		000</td <td></td> <td><0</td> <td></td>		<0	
75	Grid 175																300		300		<1600		< D	 -
176	Grid 176										<u> </u>						400		400		<1000	_	<u> </u>	 -
177	Grid 177																400		400		< 1000	_ _	2D	
178	Grid 178																400		400		<1000		∠ D	
179	Grid 179															7	400		400		<1000	_	<d< td=""><td></td></d<>	
180	Grid 180								.		\Rightarrow	6		2)	<u>\\\/</u>		400		400		< 1000	<u> </u>	< D	
۶۱	Grid 181									ļ		11	"				400		400		000</td <td></td> <td><u> </u></td> <td></td>		<u> </u>	
182	Grid 182										_						500		500		<1000		1 V D	
183	Orid 183									┸							500		500_		<1000		<d< td=""><td></td></d<>	
84	Grid 184															<u> </u>	550		550		< 1000		< D	·
185	Grid 185	· · · · · · · · · · · · · · · · · · ·										<u> </u>				<u> </u>	500		500		<1000		40	
186	Grid 186													-			400		400		< 1000		<d< td=""><td></td></d<>	
87	Grid 187	<u>.</u>															300		300		41000		< D	
88	Grid 188		NΑ	Λ	A	N	4	N	4	NΑ			N,	4	٨	14	1500		2000		<1000		15,000	
189	Grid 189	73" disc	C/F	71	NA	6	0.5	6/	A	5/1			23.9	N	3	70.5	3000		25,000		<1000		71,000,000	
190	Grid 190		NA	<u> </u>	A	N.		N/		ΝA	N	Ά	N	A	N	/ <u>A</u>	1000	NA	1000	NA	<1000	NA	4D	NA

BD-6002-601R (12/98)

SNF	PROJECT RADIOL	OGICAL SUI	RVEY F	REPO	RT - (Conti	nuatiç	n Shee	t - Dat	a Sun	ey Repo	nt No. 00 5 1 1		.	Page	10 of	16
. [Note	DOS!	E RATE eld (≥ 3	MEASI	JREMEN C = Conta	rs ct (<u><</u> 1 cm)			. · · · ·		AMINATIC	ON MEASUR	EMENT	5	,
Na.	Description	Dist. Note 1	WO mR/hr	WC mR/hr	CF ₆	CF,	Neutro Dose	Dose	Dose	· I	Bkg cpm		Direct n/PA	Remov dpm/10	rable 0 cm ²	To dpm/10	lal XV cm²
		cm				<u> </u>	mrem/h	r mrem/h	mrem/	hrβy	α	ργ	a	βγ	α	βγ	a
91 (Grid 191	NA	NA	NA	NA	NA	NĄ	NA	NA	450	NA	450	NA	<u>≺/∞0</u>	NA	< D	NA
92 (Grid 192									35		350		000</td <td></td> <td>< D</td> <td></td>		< D	
93 (Grid 193									40		400		<1000		< D	
94 (<u>3rid</u> 194									40		4.00				40	
95 @	Grid 195									35		350		000</td <td></td> <td>< D :</td> <td></td>		< D :	
36 (3rid 196									30	0	300		<1000		<0	
97 G	inid 197									300		300		< 1000		<0	
97 6	irid 198									300		300_		< 1000		∠D_	
99 6	Srid 199									300		300		< 1000		ZD	
10 0	grid 200									30		300		4/000		ZD.	
01 6	Srid 201	·			·					30		300	7.	<1000		< D:	
02 6	frid 202			.*						30		300		000</td <td></td> <td>< D.</td> <td></td>		< D.	
03 6	Frid 203									300		300		< 1000		< D	
04 6	Brid 204						·			30		300		<1000		<d_< td=""><td></td></d_<>	
<u> </u>	ini d 205									300		300	<u> </u>	000</td <td></td> <td><<u>D</u></td> <td></td>		< <u>D</u>	
06 G	frid 206					-	5 /2			300		300		<1000		</td <td></td>	
07 G	rid 207					<i>NI</i> =		•		30		300		<1000		40	
08 G	rid 208				V	74	4			30		300		<i∞0< td=""><td></td><td>< D</td><td></td></i∞0<>		< D	
9 0	Srid 209			;						300) _ N	300		<1000		< D	
0 6	orid 2/0									30	300	-300		< 1000		< D	
11 G	irid 211									30	0 309	300		< 1000		< D	
12 G	srid 212											1000		41000		30,000	
	srid 213											NA .				NA	
	3nd 214											5000		4.1000	•	210,000	
	irid 215	NA	NΑ	NA	NΑ	NA	NÅ	NA	NA			1000	ΝÀ	<1000	MA	4 D	NA

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SN	F PROJECT RADIOLOGIC	CAL SUI	RVEY	REPO	RT.	· C	ont	inua	tion	Sheet	- D	ata	Surve	Rep	NO. KOOO	511		Page	11_or	<u> d </u>
			No	DOS	E RA` ield (≥	TE M 30 c	EAS	UREM C = Co	ENTS intact	(≤ 1 cm)		•			CONT	AMINATIO	ON MEASUF	REMENT	5	
No.	Description	Dist. Note 1	WO mR/hr	WC mR/hr	CF		CF,	Do	utron ose	Shallow Dose		eep lose	BI			Direct n/PA	Remo dpm/10	vable 00 cm ²	To dpm/10	ital 00 cm²
		cm	, , , ,			<u>ا</u>		mre	m/hr	mrem/hr	mr	em/h <i>t</i>	βγ	α	ργ	α	βγ	α	βγ	α
216	Grid 216	NA	ŅΑ	NA	N	4	Nγ	<u> </u>	A.	NA	_	NA	300	NA		NA	< 1000	MA	4D	Ν4
217	Grid 217			<u> </u>		_	\perp	<u> </u>	1	<u> </u>	<u> </u>		300		300		00</td <td></td> <td>< D</td> <td></td>		< D	
218	Grid 218			 									300		300		<1000		< D	
49	Grid 219 .			<u> </u>			_	<u> </u>			<u> </u>	<u> </u>	300	\perp	300	•	<1000		∠D_	
220	Grid 220				1_1	\perp	1	<u> </u>			<u> </u>		300		300		<100C		< D	
221	Grid ZVI	_ _ :				_					L	_	300		300	<u> </u>	<1000	·	< 0	
222	Grid 222		•			_	_	ļ_			Section .	ىدرى <u>د.</u>	300	\bot	300		< 1000		∠ D-	
223	Grid 223						\perp	_					300	_	300		K1000		<u> </u>	
224	Grid 224					_ _							300	\bot	300		< 1600		<d_< td=""><td></td></d_<>	
225	Grid 225							<u> </u>				<u> </u>	300		300		<1000		40	
224	Grid 226					\perp		<u> </u>				<u> </u>	500		500		<u> </u> ≺2000	<u> </u>	40	_
127	Grid 227		·			_	1				<u> </u>	_	500		≤ 80	<u> </u>	0∞</td <td></td> <td>≺D_</td> <td></td>		≺D_	
228	Grid 228					_							500	\perp	500		< 1000	<u> </u>	< D	
229	Grid 225					1			<u> . </u>				500		500		< 1000		<u>≺D</u>	
230	Grid 230					_ _					<u> </u>		500	_	500		K1000	<u> </u>	< D	
23/	Grid 231					۷,		\[F	_	<u> </u>		<u> </u>	500	_	500		<1000	<u> </u>	<d< td=""><td>· `</td></d<>	· `
382	Grid 232						(12	M.			500		500		K1000		< D :	
233	Grid 233					-1	1:	<u> </u>			Ŀ		500		500_		<1000		< D	
234	arid 234										·		500		500		<1000		<u><1)</u>	
235	Grid 235							<u> </u>					500		500		41000		<0	
236	Grid 236					F							500		800		<1000		18,000	
237	Grid 237												500		500		< 1000		< D	
238	Grid 238												50,0		2000		<1000		90,000	
239	Grid 239												300		1500		41000		72,000	
240	Grid 240	NA.	ΝΆ	NA	N	$+\Gamma$	NA	N	A	WA	٨	VA.	300	MA	3∞	NA	<1000	NΑ	< D	NA

BD-6002-601R (12/98)

SN	F PROJECT RADIOLOGIC	AL SI	JŖV	EY	REP	OR	T - (Cont	inu	atior	Sheet	- L	ata	Surve	Repo	K000	<u> 511</u>		Page	12 01	jb:
	•			Note	1 F =	OSE Flei	RATE ld (≥ 30	MEAS	URE C = 0	MENTS Conlact	(<u><</u> 1 cm)			ļ 		I. CONT	OITANIMA	N MEASU	REMENTS	S	
No.	Description	Dist. Nate		NO R/hr	WC mR/		CF _B	CF		eutron Oose	Shallow Dose	1 (Deep Dose		kg m		Direct 1/PA	Remo dpm/10	vable 00 cm ²	dpm/10	tal 00 cm ²
		cm							m	rem/hr	mrem/hr		tem/pt	βγ	α	βγ	α	βγ	α	βγ	α
241	Grid 241	· NA		JA.	N	A.	NA	1 1/	<u> </u>	NA	NA	_^	JA.	300	NA	300	NA	<1000	NA	<d< th=""><th>NA</th></d<>	NA
42	Grid 242		_ _	_		_		1	-	-	-	<u> </u>		300		300		<1000	 	< D	
43	Grid 243						\perp			ļ		_		300		300		<1000		< D	 -
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45	Grid 245									_		_	<u>. </u>	300		300		₹1000		4P	
46	Grid 246								<u> </u>	<u> </u>		$oxed{}$		300		300		∠1000		< D	
47	Grid 247					_			丄	<u> </u>		L		300		300		< 1000		<u> </u>	
48	Grid 248											L	Ш	300		300		< 1000		< D	
49	Grid 249				.				1_	'		_		300		300		<1000		< D	 -
50	Grid 250 .		Ш.	1					⊥_					300		300		<1000		<0	
51	Grid 251			_					_			<u> </u>		300		300		< 1000		< D	- -
52	Grd 252	1_1		1_								L	<u> </u>	300		300		<1000		<d< td=""><td> -</td></d<>	-
53	Grid 253			<u> </u>		_			_			L		300		300		<1000	<u> </u>	< <u>0</u>	
54	Grid 254											_		300	<u> </u>	300		<1000		(1)	-
<u>55</u>	Grid 254 255	- ·	1	↓_					<u> </u>	_		_		300		300		4/000		<u><</u> D	_ -
50	Gid . 256			_					\bot		7	L		300		300		< 1000	 	1 < D	-
<u>57</u>	Grid: 257			<u> </u>		_	_/-	1	4			<u> </u>		300		300		< 1000		1 LD	
58	Grid 258					_;	_(_	1]))	1	n	Ŀ		360		300		21000		4D	-
<u>59</u>	Girl 259		\perp			7		2	1			_		300		300	<u> </u>	< 1000		< D	
60	Gid 260		:						<u> </u>	<u> • </u>				300		300	 	<1000 ∠1000		\ \L	
61	Grid 261											L		300		600		< 1000		18,000	
62	Grid 262:			<u> </u>					_			_		300		700		<1000		24,000	1
43	Grid 263.			_			\perp		_			$oxed{oxed}$		500		1000		< 1000		30,000	
	Grid 264													500		500		<1000		YD_	
165	Grid 265	NA	A	A	N,	4	NA	NA	-	VA	NA		NA	500	NA	500	NA	<1000		<u> < D</u>	1 NA

BD-6002-601R (12/98)

SN	F PROJE	CT RADIOLOG	SICAL S	SUF	RVEY	RE	PC	RT	- C	ont	inu	atio	n Sh	eet	- D	ata	Survey		n No. K 000.	511		Pag	e <u>//</u> o	1_16
					N	ole 1	DO: F = F	SE R	ATE (≥30	MEAS cm)	UREI C = C	MENT ontac	S l(<u>≤</u> 10	m)					CON	TAMINATI	ON MEASU	REMEN	rs	
No.		Description	Di No	te 1	WO mR/n	, ,	WC nR/h/		CF _B	CF	1 (eutron Oose	D)	allow ose	D	eep ose	g) cp			Direct n/PA	Rem dpm/1	ovable 00 cm2	dpm/1	olal 100 cm ²
								1-	<u>,</u>		<u> </u>	em/hr	1771	m/hr	II H C	m/hr	рγ	α	βr	a	βγ	α	ργ	a
291	<u>Grid</u>	291	1 ~	<u>A</u>	NA	_ 1	<u>VA</u>	4	WA-	NA	11	JA.	1	JA	^	JA	300	NΑ	300	NA	< 1000	NA		NA
292	<u>6-10</u>	292				-\-			4_	\perp	_	4	_	↓	<u> </u>	_	250	_	250		1000		< D	
53	Grid	293	·	\square		_ _		+	+		_	- -	_			 	250		250		< 1000		<u> ≺⊅</u>	-
941	<u>Grid</u>	294				4_					1-		 				250		250		<u> <1000</u>		(<)	
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96	Grid .	296							+	_	-	4	 				250	_	250	 -	< 1000		< D	 -
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9	Grid	299				- -	-	┷	11		_	 	ļ				.300		300		< 1000		<u> </u>	
00	Grid	300				_ _	<u> </u>	丄			1_	<u> </u>					250	_	250		<1000		< 0	
	Grid	301						1_	1.1		<u> </u>	1					300		300		<u> <1000</u>		$\angle D$	-
22	Grid	302						1_	77.		L						300		300		< 1000		1.YD	
23	Grid	303					<u> </u>	\perp									300		300		< 1000		140	
04	Grid	304															300		300		< 1000		LYD	
8	Grid	305															300		300		4,000	·	<d< td=""><td></td></d<>	
06	Grid	306						\perp									300		300		<1000		ZD	
01	Grid	307					_	\perp			<u> </u>		7				300		300		K1000		< D	
28	Grid	308				- _				\Rightarrow			<u>Y</u>				300		300		21000		<0	
29	Grid	309						(C			1		b				300		300		K 1000		イワ	
0	Grid	3/0						1	2								300		300		< 1000		<d< td=""><td></td></d<>	
ji i	Grid	311.		.1												}	300		300		< 1000		<0	
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	Grid	314				T		T	\prod								250		250		4/000		415	
15	Grid	315	N	4	NA	1-	ÚΑ	1	A	NA	1	A	W	4	Λ/	A	250	NA	 	NA	<1000	 - - - - - - - - - -	125	N

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BD-6002-602R (12/98) W \sqrt{N} MY AM AN W MY AN AN AN AN AM. YN AW \sqrt{N} Rev. AN AN MY AN AU 3' Perimate Around Rod Pad クラ 000/> 05% 520 050 Grid 578 (J z 0001 > 057 00017 125 11.19 050 QST ME σZ 900/2 057 052 Grid 373 (17 00017 052 222 (म! प 225 (17 052 ME 9S7 المناط 125 0001 > 057 7> 350 00017 QS7 052 975 Bird (7) 052 48 b vo 418 000/> 052 J> 001> 952 X1E bira 318 05Z 07 0001> 057 052 [15 bine LIE マウ YN ∇N ₩¥. \sqrt{N} 队为 916 AN 00/2 \overline{AM} -052 AN AN ΔN d> 520 AN AMAU ΥQ λd ΥŲ ĸ n mrembr mremyhr mrem/hr шэ wgwt ώςγνι ΜΟ c_{L} Shallow Neutron Dose Dist. PaloN Deep Smo 001/mgb Removable 2m2 ODI\mqb Gross Direct Cpm/PA шдэ BKG Description ON. Mole f = Fleid (2.30 cm) C = Contact (5.1 cm)CONTAMINATION MEASUREMENTS KOOOZII Page 15 of 18 SNF PROJECT RADIOLOGICAL SURVEY REPORT - Continuation Sheet - Map/Sketch Survey Report No.

SNF	PRO.	PROJECT RADIOLOGICAL SURVEY REPORT - Continuation Sheet - Map/Sketch													Survey Report No. K000511								e <u> 6</u> of	18			
	· ·							me	metal Map/Sketch								chil	te A									
		2	3	4	5	6	7	В	2	JU		12	13	14	15	01	17	18	/9	20	21	- 22	123	24	- II - F		
. 1	26	27	2.8	27	30	3/	32	33	34	36	36	37	38	35	40	41	42	43	44	45	112	47		9 <u>4</u>	7 30		
	S	52	53	3/	-35	52	57	78	54	60	6,1	62	63	64	45	64	61	∠ B	11/1	70	1/2		2 7	3	75		
FCA/RA	76	7	76	79	<u>80</u>	<u>SL</u>	82	63	84	85	86	87	28	89	50	2	92	В	94	85	90	9	1 9	<u>ย</u>	7 100		Ca
	101	27	103	101	10.5	1416	/07	108	107	110	111	112	<i>"3</i>	fell	115	110	117	118	119	120	121	/22		; <u> </u> /2	125	,}	
	126	127	128	129	70	134	132	133	131/	135	134	137	138	137	140	141	142	143	144	M5	710	147		ع إح	M. 150	 RA/FC	
المرام	<u>i51</u>	152	153	134	155	152	157	/28	<u>157</u>	100	101	102	n3	إيسال	1105	100	147	168	169	טרנ	17/	/72	///		175		7
	776	177	128	/75	180	181	IPZ	163	184	18;	150	157	1.98	161	150	171	192	123	124	F15"	M	177	198	199	24,0		
	201	202	205	201	205	200	207	205	202	טעיג	211	2/2	213	214	215	214	217	215	219	است	ريج	22	223	224	235		
	226	277	<i>178</i>	227	230	23/	232	233	234	22	214	237	<u>26 </u>	239	240	<u>241</u>	212	27.2	244	245	146	247	246	244	252) .		
	251	352	253	234	255	252	257	258	2551	260	241	24.2	2107	المعتثم	215	24	267	268	201	270	27,	272	223	274	27:5		
	274	277	278	271	280	281	202	283	201	265	296	257	268	287	290	AII.	292	283	25/	255	290	2,7	248	259	300		
	301	302	305	<i>3</i> ⇔	305	906	307	<i>3</i> €8	30%	310	3/1	3/2	313	إساق	315		<i>317</i>	3,8	319	320	32/	322	323	بعتق ا	325	_	
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11	J				•		• •														•			•			